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(54) Title: METHODS FOR PREPARING PURIFIED LIPOPEPTIDES

(57) Abstract: The present invention relates to crystalline and crystal-like forms of lipopeptides, including daptomycin, a lipopeptide antibiotic with potent bactericidal activity against gram-positive bacteria, including strains that are resistant to conventional antibiotics. The present invention relates to methods of purifying lipopeptides, including daptomycin, a lipopeptide antibiotic with potent bactericidal activity against gram-positive bacteria, including strains that are resistant to conventional antibiotics. The present invention also relates to pharmaceutical compositions comprising the purified form of the lipopeptide and methods of using these compositions.

WO 02/059145 A1

## METHODS FOR PREPARING PURIFIED LIPOPEPTIDES

### CROSS-REFERENCE TO RELATED APPLICATIONS

5

The present application claims the benefit of United States Provisional Application Number 60/256,268, filed December 18, 2000; Serial Number 60/274,741, filed March 9, 2001; Serial Number \_\_\_\_\_ filed December 13, 2001; and Serial Number \_\_\_\_\_ filed December 13, 2001, the contents of which are incorporated  
10 herein by reference.

### TECHNICAL FIELD OF THE INVENTION

The present invention relates to crystalline and crystalline-like forms of lipopeptides, including daptomycin, a lipopeptide antibiotic with potent bactericidal  
15 activity against gram-positive bacteria, including strains that are resistant to conventional antibiotics. The present invention also relates to processes for preparing crystalline or crystal-like forms of the lipopeptide and to methods of purifying lipopeptides including daptomycin. The present invention also relates to pharmaceutical compositions comprising the purified form of the lipopeptide and methods of using these compositions.

20

### BACKGROUND OF THE INVENTION

The rapid increase in the incidence of gram-positive infections—including those caused by antibiotic-resistant bacteria—has sparked renewed interest in the development of novel classes of antibiotics. One such class is the lipopeptide antibiotics, which  
25 includes daptomycin. Daptomycin has potent bactericidal activity *in vitro* against clinically relevant gram-positive bacteria that cause serious and life-threatening diseases. These bacteria include, but are not limited to, resistant pathogens, such as vancomycin-resistant enterococci (VRE), methicillin-resistant *Staphylococcus aureus* (MRSA), glycopeptide intermediary susceptible *Staphylococcus aureus* (GISA), coagulase-negative  
30 staphylococci (CNS), and penicillin-resistant *Streptococcus pneumoniae* (PRSP), for which there are very few therapeutic alternatives. See, *e.g.*, Tally et al., 1999, Exp. Opin. Invest. Drugs 8:1223-1238. Daptomycin's inhibitory effect is a rapid, concentration-

dependent bactericidal effect *in vitro* and *in vivo*, and a relatively prolonged concentration-dependent post-antibiotic effect *in vivo*.

Daptomycin is described by Baltz in Biotechnology of Antibiotics, 2nd Ed., ed. W.R. Strohl (New York: Marcel Dekker, Inc.), 1997, pp. 415-435. Daptomycin, also  
5 known as LY 146032, is a cyclic lipopeptide antibiotic that can be derived from the fermentation of *Streptomyces roseosporus*. Daptomycin is a member of the factor A-21978C<sub>0</sub> type antibiotics of *S. roseosporus* and is comprised of a decanoyl side chain linked to the N-terminal tryptophan of a cyclic 13-amino acid peptide (Fig. 1).  
Daptomycin has an excellent profile of activity because it is highly effective against most  
10 gram-positive bacteria; it is highly bactericidal and fast-acting; it has a low resistance rate and is effective against antibiotic-resistant organisms. The compound is currently being developed in a variety of formulations to treat serious infections caused by bacteria, including, but not limited to, methicillin-resistant *Staphylococcus aureus* (MRSA) and vancomycin-resistant enterococci (VRE).

15 A number of United States Patents describe A-21978C<sub>0</sub> antibiotics and daptomycin-related lipopeptides including daptomycin (LY 146032). These patents also describe methods of producing and isolating the A-21978C<sub>0</sub> antibiotics and daptomycin-related lipopeptides.

United States Patents RE32,333, RE32,455, 4,800,157, 4,874,843, and 4,885,243  
20 describe methods of synthesizing and isolating daptomycin from fermentation cultures of *Streptomyces roseosporus*. United States Patents RE32,310, RE32,311, 4,537,717, 4,482,487 and 4,524,135 describe A-21978C<sub>0</sub> antibiotics and methods of deacylating the A-21978C<sub>0</sub> antibiotic and reacylating the peptide nucleus and antibiotic derivatives made by this process. United States Patent 5,912,226 (hereafter the '226 patent) describes the  
25 identification and isolation of two impurities produced during the manufacture of daptomycin, anhydro-daptomycin and the  $\beta$ -isomer form of daptomycin. None of these United States patents discloses a method for precipitating or crystallizing a lipopeptide in a manner to increase purity of the lipopeptide.

United States Patent 4,439,425 (hereafter the '425 patent) discloses a crystalline  
30 lipopeptide and a method of crystallizing the lipopeptide. The lipopeptide disclosed in the '425 patent is structurally dissimilar from daptomycin and daptomycin-related lipopeptides. United States Patent 5,336,756 (hereafter the '756 patent) also discloses a crystalline cyclic lipopeptide comprising a hexapeptide. The crystalline cyclic

lipopeptide disclosed in the '756 patent is also structurally dissimilar from daptomycin and daptomycin-related lipopeptides. The '756 patent discloses that the lipopeptide, an echinocandin-type compound, can be obtained when aqueous n-propanol is employed as the crystallizing solvent. See, e.g., cols. 1-2 of the '756 patent. Neither the '425 patent  
5 nor the '756 patent disclose methods of crystallizing or precipitating daptomycin or a daptomycin-related lipopeptide, nor do they disclose methods of crystallizing or precipitating lipopeptides produced by *Streptomyces*.

It would be advantageous to develop a method of crystallizing or precipitating daptomycin and daptomycin-related lipopeptides to provide an improved purification  
10 method for these lipopeptides. In addition, a crystalline or highly purified precipitated form of daptomycin or other daptomycin-related lipopeptide would be useful in formulating pharmaceutical compositions for treating bacterial infections. Further, a crystalline or highly purified precipitated form of daptomycin or daptomycin-related lipopeptide would be useful in a method to make a sterile product, particularly bulk sterile  
15 product. Thus, there is a need for methods to produce crystalline or precipitated daptomycin and daptomycin-related lipopeptides and the crystalline or precipitated forms of the lipopeptides produced thereby. However, there has been no simple and robust method that has been effective in crystallizing or precipitating daptomycin or a daptomycin-related lipopeptide that results in a lipopeptide that is more pure after  
20 crystallization or precipitation than before.

#### SUMMARY OF THE INVENTION

The instant invention addresses these problems by providing crystalline and crystalline-like forms of lipopeptides, particularly daptomycin and daptomycin-related  
25 lipopeptides and methods for producing them. In one embodiment, the invention provides methods for crystallizing lipopeptides. In another embodiment, the methods provide a lipopeptide that is more pure after crystallization or precipitation than before crystallization or precipitation.

The invention also provides robust processes for producing and purifying  
30 lipopeptides comprising, *inter alia*, crystallizing or precipitating lipopeptides. In one embodiment, the crystallizing or precipitating steps of the processes are used to purify the lipopeptides. In another embodiment, the processes are used for large-scale and/or commercial production of lipopeptides, preferably daptomycin.

The invention further provides highly purified crystalline or crystal-like forms of daptomycin and daptomycin-related lipopeptides. In one embodiment, the crystalline or crystal-like forms of the lipopeptides may be used in pharmaceutical compositions. In another embodiment, the invention comprises methods of using the pharmaceutical compositions.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 shows the structure of daptomycin.

Fig. 2 shows a photomicrograph of urchin-like crystal or crystal-like particle of daptomycin produced by the method described in Example 12.

Fig. 3 shows a photomicrograph of needle-like crystals of daptomycin.

Fig. 4 shows a photomicrograph of rod-like crystals of daptomycin.

Fig. 5 shows photomicrographs of daptomycin samples at 100X magnification. Photomicrographs of amorphous daptomycin are shown using plane transmitted light (A) and using crossed polarized light (B). Photomicrographs of daptomycin crystals are shown using plane transmitted light (C and E) and using crossed polarized light (D and F). The daptomycin crystals were produced by the protocol disclosed in Example 7.

Fig. 6 shows an x-ray powder diffraction pattern for amorphous daptomycin.

Fig. 7 shows an x-ray powder diffraction pattern for a daptomycin crystal produced by the protocol described in Example 7.

Fig. 8 shows an x-ray powder diffraction pattern for a second sample of a daptomycin crystal produced by the protocol described in Example 7.

Fig. 9 shows birefringence of a crystal-like particle of daptomycin when exposed to polarized light. The crystal-like particle was produced by the method described in Example 12.

Fig 10 shows a flow chart of an exemplary method for crystallization.

Fig. 11 shows a flow chart of an exemplary manufacturing method that does not use crystallization or precipitation. The manufacturing method uses bacterial fermentation to produce a fermentation culture containing daptomycin, and then purification of daptomycin using microfiltration, anion exchange chromatography, size exclusion ultrafiltration, hydrophobic interaction chromatography, anion exchange chromatography for solvent removal, ultrafiltration for pyrogen removal, reverse osmosis and filling vials with daptomycin. See, e.g., International PCT Publication WO 01/44274,

published June 21, 2001, herein incorporated by reference for a detailed description of this type of method.

Fig. 12 shows a flow chart of an exemplary manufacturing method of a lipopeptide compound comprising the steps of fermentation, microfiltration, anion  
5 exchange chromatography, size exclusion ultrafiltration, crystallization or precipitation, crystal or precipitate drying, and dry filling of vials with the compound. See, e.g., Example 13.

Fig. 13 shows a flow chart of an exemplary manufacturing method of a lipopeptide compound comprising the steps of fermentation, microfiltration, anion  
10 exchange chromatography, crystallization or precipitation, crystal or precipitate drying, and dry filling of vials with the compound. See, e.g., Example 14.

Fig. 14 shows a flow chart of an exemplary manufacturing method of a lipopeptide compound comprising the steps of fermentation, microfiltration, size  
15 exclusion ultrafiltration, crystallization or precipitation, crystal or precipitate drying, and dry filling of vials with the compound. See, e.g., Example 15.

Fig. 15 shows a flow chart of an exemplary manufacturing method of a lipopeptide compound comprising the steps of fermentation, microfiltration, crystallization or precipitation, crystal or precipitate drying, and dry filling of vials with  
the compound. See, e.g., Example 16.

20 Fig. 16 depicts the structure of CB-131547, a cyclic lipopeptide analog of daptomycin

## DETAILED DESCRIPTION OF THE INVENTION

### Objects of the Invention

25 One object of the present invention is to provide methods for crystallizing or precipitating lipopeptides. In one embodiment, the methods are used to crystallize or precipitate daptomycin or a daptomycin-related lipopeptide. In another embodiment, the methods increase the purity of the lipopeptide compared to the purity of the lipopeptide prior to crystallization or precipitation. The methods comprise the steps of providing an  
30 amorphous preparation of a lipopeptide and crystallizing or precipitating the lipopeptide under conditions in which the crystalline or precipitated, crystal-like lipopeptide is more pure than the amorphous preparation of the lipopeptide. In one embodiment, the amorphous preparation is no greater than 92% pure and the crystalline or crystal-like

lipopeptide purified therefrom is at least 95% pure, and may be at least 96%, 97% or 98% or more pure. In another embodiment, the amorphous preparation is no greater than 80% pure and the crystalline or crystal-like lipopeptide purified therefrom is at least 95% pure, and may be at least 96%, 97% or 98% or more pure. In another embodiment, the  
5 amorphous preparation is no greater than 60% pure and the crystalline or crystal-like lipopeptide purified therefrom is at least 95% pure, and may be at least 96%, 97% or 98% or more pure. In yet another embodiment, the amorphous preparation is no greater than 40% pure and the crystalline or crystal-like lipopeptide purified therefrom is at least 95% pure, and may be at least 96%, 97% or 98% or more pure. In another embodiment, the  
10 amorphous preparation is no greater than 20% pure and the crystalline or crystal-like lipopeptide purified therefrom is at least 95% pure, and may be at least 96%, 97% or 98% or more pure. In a further preferred embodiment, the amorphous preparation is no greater than 10% pure and the crystalline or crystal-like lipopeptide purified therefrom is at least 95% pure, and may be at least 96%, 97% or 98% or more pure.

15 Another object of the invention is to provide processes for making and purifying a lipopeptide comprising, *inter alia*, crystallizing or precipitating the lipopeptides. In one embodiment, the crystallizing or precipitating steps are used to purify the lipopeptides. In a preferred embodiment, the crystallization or precipitation is performed by batch crystallized or precipitation. In another embodiment, the process is a large-scale process  
20 for commercial production of a lipopeptide, preferably daptomycin or a daptomycin-related lipopeptide. In one embodiment, the lipopeptide is produced by fermentation. The fermentation product is then purified by a variety of purification techniques including crystallization or precipitation. In one embodiment, the crystallization or precipitation step may be used in combination with other purification techniques including  
25 microfiltration, size exclusion ultrafiltration and/or anion exchange chromatography. In one embodiment, the crystallization or precipitation step is used to replace one or more purification techniques that is used in a purification process that does not use crystallization or precipitation. In another embodiment, the crystallization or precipitation step is used to increase purification compared to the other steps without the  
30 crystallization or precipitation step. In a preferred embodiment, the method comprises a step of collecting the crystalline or crystal-like lipopeptide after crystallization or precipitation.



Another object of the present invention is to provide highly purified, e.g. sterile, crystalline or crystal-like forms of lipopeptides. In one embodiment, the lipopeptides are daptomycin or a daptomycin-related lipopeptide. The crystalline or crystal-like form of the lipopeptide may have any crystalline or crystal-like shape including urchin-like (cluster of needles joined together to visually resemble a sea urchin)(see Fig. 2), needle-like (see Fig. 3), rod-like (see Fig. 4), plate-like or flake-like. In one embodiment, the crystalline or crystal-like lipopeptide has a purity of at least 80%, and may be at least 85%, 90% pure. In another embodiment, the crystalline or crystal-like form of the lipopeptide has a purity of at least 95%, and may be at least 96%, 97%, 98% pure or more.

A further object of the present invention is to provide a pharmaceutical composition comprising a crystalline or crystal-like form of a lipopeptide. In one embodiment, the lipopeptide is daptomycin or a daptomycin-related lipopeptide. In one embodiment, the pharmaceutical comp. is enterically coated for oral administration or is formulated in the form of micronized particles or microspheres. In other embodiments, the invention provides methods for administering the pharmaceutical compositions to subjects in need thereof.

#### Definitions

Unless otherwise defined, all technical and scientific terms used herein have the meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. The practice of the present invention employs, unless otherwise indicated, conventional techniques of chemistry, biochemistry, biophysics and microbiology and basic terminology used therein.

The term "lipopeptide" refers to a molecule that comprises a lipid-like moiety covalently linked to a peptide moiety, as well as salts, esters, amides and ethers thereof. The term "lipopeptide" also encompasses protected forms of lipopeptides in which one or more amino, carboxylate or hydroxyl groups are protected. See, e.g., "Protective Groups in Organic Synthesis" by Theodora W. Greene, John Wiley and Sons, New York, 1981 for examples of protecting groups. In one embodiment, the lipopeptide is an antibiotic. In another embodiment, the lipopeptide is LY 303366, echinocandins, pneumocandins, aculeacins, viscosin, surfactin, plipastatin B1, amphomycin or the lipopeptide derivative disclosed in United States Patent 5,629,288. These lipopeptides are known in the art.

See, e.g., United States Patent 5,202,309 and International PCT Application WO 00/08197. In another embodiment, the lipopeptide is a daptomycin-related molecule. In another embodiment, the lipopeptide is daptomycin.

A “daptomycin-related molecule” includes, *inter alia*, daptomycin, A54145 or  
5 other lipopeptide that is structurally related to daptomycin, such as a daptomycin-related lipopeptide, including all stereoisomers that may be made at any chiral centers present in these molecules.

A “daptomycin-related lipopeptide” includes, without limitation, a lipopeptide disclosed in United States Patent 4,537,717, 4,482,487, RE32,311, RE32,310, and  
10 5,912,226, currently in reissue as United States Application No. 09/547,357. Daptomycin-related lipopeptides also include those disclosed in International PCT Publication WO 01/44272, published June 21, 2001; International PCT Publication WO 01/44274, published June 21, 2001; and International PCT Publication WO 01/44271, published June 21, 2001; all of these applications are specifically incorporated herein by  
15 reference. The daptomycin-related lipopeptides disclosed in the above-identified applications relate to synthetic and semisynthetic lipopeptides in which the ornithine and/or kynurine residues, and/or the fatty acid side chain of daptomycin, are modified. Daptomycin-related lipopeptides further include an A-21978C<sub>0</sub> antibiotic in which the n-decanoyl fatty acid side chain of daptomycin is replaced by a n-octanoyl, n-nonanoyl, n-undecanoyl, n-dodecanoyl, n-tridecanoyl or n-tetradecanoyl fatty acid side chain.  
20

The term “daptomycin” refers to the n-decanoyl derivative of the factor A-21978C<sub>0</sub>-type antibiotic that contains an  $\alpha$ -aspartyl group. “Daptomycin” is synonymous with LY 146032.

The term “anhydro-daptomycin” refers to a daptomycin-related lipopeptide in  
25 which an  $\alpha$ -aspartyl group of daptomycin is cyclized to a succinimido group. See, e.g., the ‘226 patent for the structure of anhydro-daptomycin.

The term “ $\beta$ -isomer” or “ $\beta$ -isomer of daptomycin” refers to a daptomycin-related lipopeptide that contains a  $\beta$ -aspartyl group instead of an  $\alpha$ -aspartyl group. See, e.g., the ‘226 patent for the structure of  $\beta$ -isomer of daptomycin.

30 The term “isolated” refers to a compound or product that is at least 5%, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80% or 90% of the compound present in a mixture. It will be understood that the term “isolated” also refers to a compound that is at least 5-

10%, 10-20%, 20-30%, 30-40%, 40-50%, 50-60%, 60-70%, 70-80% or 80-90% of the compound present in the mixture group. The percentage of compound in a mixture may be measured by any means known in the art, as described below for measuring purity of a compound.

5           “Substantially pure” refers to a sample having at least 95% of a desired compound. Preferably, daptomycin is “substantially pure” when at least 95% to at least 97% of a sample is daptomycin. Similarly, a daptomycin-related lipopeptide, is “substantially pure” when at least 95% to at least 97% of a sample is a daptomycin-related lipopeptide.

10           Daptomycin or a daptomycin-related lipopeptide is “essentially pure” when at least 98% to at least 99% of a sample is daptomycin or a daptomycin-related lipopeptide, respectively.

            Daptomycin or a daptomycin-related lipopeptide is “substantially free” of another compound when the other compound is present in an amount that is no more than 1% of  
15           the amount of the daptomycin or the daptomycin-related lipopeptide preparation, respectively.

            Daptomycin or a daptomycin-related lipopeptide is “essentially free” of another compound when the other compound is present in an amount that is no more than 0.5% of the amount of the daptomycin or the daptomycin-related lipopeptide preparation,  
20           respectively.

            Daptomycin or a daptomycin-related lipopeptide is “free” of another compound when the other compound is present in an amount that is no more than 0.1% of the amount of the daptomycin or the daptomycin-related lipopeptide preparation, respectively. Alternatively, daptomycin or a daptomycin-related lipopeptide is “free” of  
25           another compound when the compound cannot be detected by HPLC under conditions of maximum sensitivity in which a limit of detection is approximately 0.05% or less of the amount of the daptomycin or the daptomycin-related lipopeptide preparation, respectively.

            “Purified” daptomycin refers to substantially pure daptomycin, essentially pure  
30           daptomycin, or a salt thereof, or to daptomycin or a salt thereof which is substantially free, essentially free, or free of another compound. Similarly, a “purified” daptomycin-related lipopeptide refers to a substantially pure daptomycin-related lipopeptide, an essentially pure daptomycin-related lipopeptide, or a salt thereof, or to a daptomycin-

related lipopeptide or a salt thereof which is substantially free, essentially free, or free of another compound.

“Crude” daptomycin refers to daptomycin or a salt thereof that is less than 90% pure. Similarly, “crude” daptomycin-related lipopeptide refers to a daptomycin-related lipopeptide or a salt thereof that is less than 90% pure.

“Semi-purified” daptomycin refers to daptomycin or a salt thereof that is at least 90% pure and less than 95% pure. Similarly, “semi-purified” daptomycin-related lipopeptide refers to a daptomycin-related lipopeptide or a salt thereof that is at least 90% pure and less than 95% pure.

The purity of daptomycin, daptomycin-related lipopeptide or of another lipopeptide refers to the lipopeptide prior to its formulation in a pharmaceutical composition. The purity of the lipopeptide is referred to by “percent purity.” The measure of purity is not a measure of degree of crystallinity of the crystalline preparation. The purity may be measured by any means including nuclear magnetic resonance (NMR), gas chromatography/mass spectroscopy (GC/MS), liquid chromatography/mass spectroscopy (LC/MS) or microbiological assays. One preferred means for measuring the purity of daptomycin is by analytical high pressure liquid chromatography (HPLC). Two methods of analytical HPLC are described in International PCT Publication WO 01/53330, published July 26, 2001, which is herein incorporated specifically by reference.

A “lipopeptide crystal” refers to one or more crystals of a lipopeptide or of a lipopeptide salt. The determination of a lipopeptide as a crystal can be determined by any means including, *inter alia*, optical microscopy, electron microscopy, x-ray powder diffraction, solid state nuclear magnetic resonance (NMR) or polarizing microscopy. Microscopy can be used to determine the crystal length, diameter, width, size and shape, as well as whether the crystal exists as a single particle or is polycrystalline.

A lipopeptide or lipopeptide particle is “crystal-like” if it is determined to have crystalline characteristics when determined by one means, e.g., visually or by optical or polarizing microscopy, but does not have crystalline characteristics when determined by another means, e.g., x-ray powder diffraction. A lipopeptide that is “crystal-like” may be crystalline under certain conditions but may become non-crystalline when subjected to other conditions.

A “crystalline lipopeptide” or a “crystalline form of a lipopeptide” refers to a preparation of a lipopeptide or salt thereof that comprises lipopeptide crystals. In one

embodiment, a crystalline lipopeptide may comprise some amount of amorphous lipopeptide. In one embodiment, the crystalline lipopeptide comprises more than 50% by weight of lipopeptide crystals. In another embodiment, the crystalline lipopeptide comprises more than 60%, 70%, 80%, 90% or 95% of lipopeptide crystals. The crystalline lipopeptide may comprise 50-60%, 60-70%, 70-80%, 80-90% or 90-95% of lipopeptide crystals. In another embodiment, the crystalline lipopeptide comprises more than 95% of lipopeptide crystals, e.g., at least 96%, 97%, 98% or 99% lipopeptide crystals or 100% lipopeptide crystals. The crystalline lipopeptide may also comprise anywhere from 95-100% lipopeptide crystals. The percent by weight of lipopeptide crystals refers to the lipopeptide preparation prior to its formulation in a pharmaceutical composition.

An "amorphous" form of a lipopeptide refers to a lipopeptide preparation that comprises few or no lipopeptide crystals or crystal-like lipopeptides (or crystal-like particles) as defined herein. In one embodiment, an amorphous lipopeptide comprises less than 20% by weight of lipopeptide crystals or crystal-like lipopeptides. In another embodiment, an amorphous lipopeptide comprises less than 10% by weight of lipopeptide crystals or crystal-like lipopeptides. In another embodiment, an amorphous lipopeptide comprises less than 5% by weight of lipopeptide crystals or crystal-like lipopeptides. In a still further preferred embodiment, an amorphous lipopeptide comprises less than 1% by weight of lipopeptide crystals or crystal-like lipopeptides.

"Batch crystallization" refers to a method in which the lipopeptide of interest is mixed with the crystallization reagents in solution and the lipopeptide is allowed to crystallize in solution. "Batch precipitation" refers to a method in which the lipopeptide is mixed with precipitation reagents in solution and the lipopeptide is allowed to precipitate in solution. In one embodiment, the crystalline or precipitated preparation is collected from the solution. In another embodiment, the crystalline or precipitated preparation is collected by filtration or centrifugation.

"Organic precipitant" refers to a polyethylene glycol (PEG) or polyethylene glycol monomethyl ether (PEG MME) or compounds that are chemically similar.

"Salts" refer to ionic compounds. These ionic compounds may act as precipitants.

"Low molecular weight alcohols" are organic compounds containing at least one alcohol functional group, and eight carbon atoms or less. For example, low molecular weight alcohols include, without limitation, methanol, isopropanol, and *tert*-butanol.

“Polyhydric alcohols” refer to compounds that contain more than one alcohol group, and less than eight carbon atoms. Polyhydric alcohols, for example, include, without limitation, 1,6 hexanediol, ethylene glycol, propylene glycol, glycerol, 1,2-propanediol, 2-methyl-2,4-pentanediol and 1,4 butanediol.

5 “Container” refers to a receptacle for holding goods. For example, a container may include, without limitation, an ampule, vial, tube, bottle, or cylinder.

#### Methods for Producing Purified Lipopeptides

One object of the invention is to provide a method for purifying a lipopeptide  
10 comprising the steps of providing an amorphous preparation of a lipopeptide and crystallizing or precipitating the lipopeptide. In one embodiment, the lipopeptide has a higher degree of purity after crystallization or precipitation than prior to being subjected to crystallization or precipitation. Lipopeptides may be crystallized by hanging drop, sitting drop or sandwich drop vapor diffusion, liquid-liquid or free interface diffusion,  
15 microdialysis or dialysis, slow solvent evaporation, sublimation, or microbatch or batch crystallization. In general, a lipopeptide may be precipitated in a similar way, preferably a lipopeptide is precipitated by batch precipitation. In a preferred embodiment, the crystallized or precipitated lipopeptide is daptomycin or a daptomycin-related lipopeptide. In a more preferred embodiment, the crystallized or precipitated lipopeptide is  
20 daptomycin.

Lipopeptides may be crystallized or precipitated following the teachings of this specification. In one embodiment, a lipopeptide can be crystallized or precipitated by providing a solution comprising a lipopeptide with a low molecular weight or polyhydric alcohol, a pH buffering agent and a salt comprising a monovalent or divalent cation and  
25 allowing precipitation or crystallization to occur, as discussed further *infra*. In another embodiment, the salt has buffering capacity such that an additional pH buffering agent does not have to be present in the solution. In another embodiment, the salt comprises a divalent cation. In a preferred embodiment, the solution provided does not include PEG or PEG-MME or chemically similar compounds. In an embodiment, the method for  
30 precipitating or crystallizing the lipopeptide generally comprises the steps of:

a) mixing the lipopeptide with a salt comprising a monovalent or divalent cation, an optional pH buffering agent and a low molecular weight or polyhydric alcohol; and

b) allowing the lipopeptides to precipitate or crystallize from the solution under the appropriate temperature conditions.

The samples may be monitored, *inter alia*, for crystal or precipitate formation by microscopic examination and the yield may be followed spectrophotometrically. In a preferred embodiment, the crystallized or precipitated lipopeptide is daptomycin or a daptomycin-related lipopeptide.

In another embodiment, the lipopeptide can be crystallized by providing a solution comprising a low molecular weight or polyhydric alcohol(s), salts and an organic precipitant as discussed further infra. In a more preferred embodiment, the crystallized lipopeptide is daptomycin. In general, for batch crystallization, the lipopeptide is dissolved in a solution and low molecular weight alcohols, salts, buffers and/or organic precipitants are added to the solution. The samples are then crystallized under the appropriate temperature conditions, with or without stirring. The samples may be monitored, *inter alia*, for crystal formation by microscopic examination and the yield may be followed spectrophotometrically.

As discussed above, the lipopeptide, preferably daptomycin or a daptomycin-related lipopeptide, is crystallized or precipitated in the presence of one or more alcohols. In a preferred embodiment, the alcohol is a low molecular weight or polyhydric alcohol. Examples of low molecular weight or polyhydric alcohols include, without limitation, methanol, isopropanol, *tert*-butanol, 1,6 hexanediol, ethylene glycol, propylene glycol, glycerol, 1,2-propanediol, 2-methyl-2,4-pentanediol and 1,4 butanediol. In a preferred embodiment, the alcohol is isopropanol, *tert*-butanol, glycerol, 1,6-hexanediol, 1,2-propanediol, 1,4-butanediol, propylene glycol and/or ethylene glycol. In a more preferred embodiment, the alcohol is isopropanol.

Salts include, *inter alia*, magnesium or sodium formate, ammonium sulfate, ammonium dihydrogen phosphate, calcium acetate, zinc acetate, tri-sodium citrate dihydrate, magnesium acetate, sodium acetate, magnesium chloride, cadmium chloride, ammonium acetate, sodium chloride and lithium sulfate. In one embodiment, the salt comprises a monovalent cation, e.g., sodium. In a preferred embodiment, the salt comprises a divalent cation. In an even more preferred embodiment, the salt comprises a calcium cation, a magnesium cation or a manganese cation. In a further preferred embodiment, the salt comprises a calcium divalent cation. In one embodiment, the salt is calcium chloride, calcium acetate, zinc acetate, sodium citrate, tri-sodium citrate

dihydrate, magnesium chloride, lithium sulfate, sodium chloride, magnesium acetate, sodium acetate or a manganese salt, such as manganese acetate or manganese chloride. In a preferred embodiment, the salt is calcium acetate. Examples of other salts that comprise a divalent cation, such as a calcium cation, are known in the art, and include, *inter alia*,  
5 those listed in the 2000 Sigma catalog, herein incorporated by reference. Without wishing to be bound to any theory, it is thought that the salt cation may neutralize the negative charges on the lipopeptide, e.g., the four carboxylic acids of daptomycin. Organic precipitants include, *inter alia*, polyethylene glycols (PEGs) that can vary in average molecular weight from between 300 and 10,000, or polyethylene glycol  
10 monomethyl ether (PEG-MME). In a preferred embodiment, the organic precipitant is PEG 300, PEG 600, PEG 2000, PEG 4000, PEG 8000 or PEG 10,000.

The lipopeptide is precipitated or crystallized from a solution that is buffered to pH 5.0 to 9.5. In one embodiment, prior to being buffered, the solution has a pH of about 1.5, 2.0 or 3.0. In one embodiment, daptomycin or a daptomycin-related lipopeptide is  
15 precipitated or crystallized from a solution of approximately pH 5.5 to approximately pH 7.5. In another embodiment, the buffer has a pH of approximately 5.9 to approximately pH 6.3. In one embodiment, the buffered solution may be obtained by using a pH buffering agent. Examples of pH buffering agents include, without limitation, Tris, phosphate, citrate, HEPES, CHES, sodium acetate or 2-morpholinoethanesulfonic acid  
20 (MES), sodium borate, sodium cacodylate, imidazole and tri-sodium citrate dihydrate. In a preferred embodiment, the salt is sodium cacodylate, sodium acetate, tri-sodium citrate dihydrate, HEPES, MES, CHES, imidazole, calcium acetate and Tris-HCl. In a more preferred embodiment, the pH buffer is calcium acetate pH 6.1, sodium acetate pH 6.1, sodium cacodylate pH 6.5, tri-sodium citrate dihydrate pH 5.6, HEPES pH 7.5, imidazole  
25 pH 8, MES pH 6.0, calcium acetate pH 6 and Tris-HCl pH 8.5. In another embodiment, the solution may be buffered by using a salt that also has buffering capacity. In a preferred embodiment, the pH buffer is calcium acetate pH 6.1.

The lipopeptide is precipitated or crystallized using hanging drop vapor diffusion from a solution containing 2 to 40% low molecular weight or polyhydric alcohol, 0.001 to  
30 0.5 M salt and 0.005 to 0.2 M pH buffering agent. In a preferred embodiment, the lipopeptide is precipitated or crystallized from a solution containing 3 to 30% low molecular weight or polyhydric alcohol, 0.01 to 0.3 M salt and 0.01 to 0.1 M pH buffering agent. In a more preferred embodiment, the lipopeptide is precipitated or



crystallized from a solution containing 5 to 20% low molecular weight or polyhydric alcohol, 0.02 to 0.1 M salt and 0.02 to 0.07 M pH buffering agent. The solution provided may or may not include polyethylene glycol (PEG) or polyethylene glycol monomethyl ether (PEG-MME).

5           The lipopeptide is precipitated or crystallized using batch crystallization from a solution containing 65 to 95% low molecular weight or polyhydric alcohol, 0.001 to 0.5 M salt and 0.001 to 0.2 M pH buffering agent. In a preferred embodiment, the lipopeptide is precipitated or crystallized from a solution containing 70 to 90% low molecular weight or polyhydric alcohol, 0.005 to 0.04 M salt and 0.005 to 0.04 M pH  
10 buffering agent. In some embodiments, the lipopeptide is crystallized from a solution which also comprises 3-8% organic precipitant. In a more preferred embodiment, the lipopeptide is precipitated or crystallized from a solution containing 80 to 85% low molecular weight or polyhydric alcohol, 0.01 to 0.03 M salt and 0.01 to 0.03 M pH buffering agent. In some embodiments, the solution further comprises about 4 to 5%  
15 organic precipitant, e.g., PEG or PEG-MME. In other embodiment, the solution provided does not include polyethylene glycol (PEG) or polyethylene glycol monomethyl ether (PEG-MME).

          The lipopeptide is precipitated or crystallized at a temperature from approximately 0°C to approximately 30°C to obtain precipitate or crystal formation, respectively. In a  
20 preferred embodiment, a lipopeptide is crystallized or precipitated at a temperature of approximately 20-30°C. In a more preferred embodiment, the mixture is crystallized or precipitated at approximately 23-28°C. In an even more preferred embodiment, the mixture is crystallized or precipitated at approximately 27°C. The mixture may be crystallized or precipitated for any time period that results in crystallization or  
25 precipitation, preferably approximately one hour to approximately two weeks. In a preferred embodiment, the mixture is stored for a period of approximately three hours to approximately 24 hours, more preferably approximately 8-18 hours.

          Lipopeptide crystals or crystal-like particles may have a shape that is, without limitation, needle-like, rod-like, urchin-like, flake-like, plate-like or clusters thereof. In  
30 one embodiment, lipopeptide crystals or crystal-like particles are urchin-like, rod-like or needle-like. The shape of the crystal or crystal-like particle may be determined, *inter alia*, by optical or electron microscopy. In another embodiment, lipopeptide crystals or crystal-like particles may be any size that is at least approximately 0.5 µm in diameter in

any one dimension. In a more preferred embodiment, lipopeptide crystals or crystal-like particle are at least 5  $\mu\text{m}$ , more preferably at least 10  $\mu\text{m}$ . In an even more preferred embodiment, the lipopeptide crystals or crystal-like particles are at least 50  $\mu\text{m}$ , more preferably at least 100  $\mu\text{m}$ . The size of the crystal may be determined by any method  
5 known to one having ordinary skill in the art. See, e.g., United States Pharmacopeia (USP), pp. 1965-67.

The properties of a crystalline or crystal-like lipopeptide may be determined by any method known to one having ordinary skill in the art. The properties that can be determined include the crystalline or crystal-like lipopeptide's size, shape, birefringence  
10 properties, powder x-ray diffraction properties, solid state NMR properties, melting temperature and stability to heat, light, humidity, and degradation. In a preferred embodiment, one having ordinary skill in the art may determine whether a lipopeptide is crystalline by powder x-ray diffraction. Powder x-ray diffraction is highly useful for determining whether a preparation is crystalline when the sample is a randomly-oriented  
15 collection of small crystals. Diffraction by a mass of randomly-oriented microcrystals produces a series of lines or rings (dependent of the detector) characteristic of the molecule studied and its structure. In a preferred embodiment, powder diffraction is measured by an Automated Powder Diffraction instrument in order to determine whether a lipopeptide is crystalline. See, e.g., Atkins et al., Physical Chemistry, pp. 710-716  
20 (1978), herein incorporated by reference for a discussion of the Debye-Scherrer method for powder diffraction. Any powder diffractometer instrument known in the art that is equipped with any detector for powder diffraction that known in the art could be used to measure the diffraction pattern.

In a preferred embodiment of the invention, a lipopeptide is crystallized or  
25 precipitated using a buffering agent between approximately pH 5.0 and 9.5, a salt and an alcohol at a temperature of approximately 24-28°C for a period of approximately three to 24 hours. In a preferred embodiment, the salt is a buffering agent and comprises a divalent cation and the alcohol is a low molecular weight alcohol, and the pH is between approximately pH 5.5 and 7.5. In an even more preferred embodiment, the salt is a  
30 calcium salt, the alcohol is isopropanol and the pH is between approximately pH 5.9 and 6.3. In embodiments where the solution includes an organic precipitant, preferably the organic precipitant is PEG 4000 or PEG 8000. In another embodiment the lipopeptide is precipitated or crystallized from a solution containing 12 to 18% glycerol, 0.3 to 0.8m

salt, 0.03 to 0.08m pH buffering agent, and 12-18% PEG 600. In a still further preferred embodiment, the lipopeptide is daptomycin or a daptomycin-related lipopeptide.

Examples 2-3 provide methods for precipitating a highly pure crystal-like daptomycin.

One having ordinary skill in the art, following the teachings of the instant specification,

5 may modify the crystallization/precipitation conditions provided in the examples to crystallize or precipitate daptomycin, daptomycin-related lipopeptides, or other lipopeptides of interest. Further, although the teachings of the instant specification describe the use of a single crystallization or precipitation step in a process for purifying a lipopeptide, one having ordinary skill in the art following the teachings of the  
10 specification may use multiple crystallization or precipitation steps in a process for purifying a lipopeptide. It may be advantageous to employ multiple rounds of crystallization or precipitation as disclosed herein in order to further increase purity of the lipopeptide.

After crystallization or precipitation, one may collect the crystalline material or  
15 crystal-like precipitate by any method known in the art. In a preferred embodiment, the crystalline material or crystal-like precipitate is collected by centrifugation or filtration. In an even more preferred embodiment, the crystalline material or crystal-like precipitate is collected by filtration because filtration is easily incorporated into a large-scale process for producing a lipopeptide. After the crystalline material or crystal-like precipitate is  
20 collected, it may be washed to remove excess crystallizing or precipitating reagents. Any wash solvent known in the art may be chosen so long as it does not appreciably dissolve the crystalline material or crystal-like precipitate. An example of a wash solvent is provided in Example 12. After the crystalline material or crystal-like precipitate is washed, it may be dried by any method known in the art. Examples of drying methods  
25 include air-drying, lyophilization (freeze-drying) or desiccation. In a preferred method, the crystalline material or crystal-like precipitate is desiccated. See, e.g., Example 12. In another embodiment, the crystalline lipopeptide's stability may be determined by its residual antibiotic activity or its degradation. The antibiotic activity may be measured in a standard agar-diffusion assay against various bacterial strains. See, e.g., Example 32 of  
30 United States Patent 4,537,717, specifically incorporated herein by reference. The amount of degradation can be measured by, *inter alia*, HPLC analysis, such as that described in International PCT Publication WO 01/53330, published July 26, 2001. In a preferred embodiment, the stability of the crystalline lipopeptide is greater than that of the

amorphous form of the lipopeptide. The stability of the crystalline lipopeptide may be determined by exposing the crystalline lipopeptide and an amorphous form thereof to heat, light, humidity, and measuring the degree of degradation of the crystalline form to that of the amorphous form.

5            Degradation of the lipopeptide may be measured by determining the biological activity of the lipopeptide or any applicable physical parameter. In one embodiment, degradation may be measured by determining a particular biological activity of a lipopeptide after it has been subjected to heat, light, humidity, changes in pH or extreme pH, and comparing it to the same biological activity of the lipopeptide prior to any tests  
10           of stability. The amount of degradation may be determined, for example, by determining the percentage of biological activity remaining after the test of stability. The percentage of remaining biological activity may be compared to that of an amorphous form of the lipopeptide that has been subjected to the same test. In one embodiment, if the lipopeptide is an antibiotic, the crystalline lipopeptide may be tested for its antibiotic  
15           activity both prior to and after a test of its stability and compared to an amorphous form that has been tested prior to and after a degradation test. In a preferred embodiment, the lipopeptide is daptomycin or a daptomycin-related lipopeptide, and the biological activity test determines the amount of antibiotic activity of the lipopeptides against gram-positive bacteria.

20           Degradation of a lipopeptide may also be measured by a physical assay. In one embodiment, degradation may be measured by determining the percentage of intact crystalline lipopeptide that remains after a test of its stability. The percentage of remaining intact lipopeptide may be compared to that of an amorphous form of the lipopeptide that has been subjected to the same test for stability. In a preferred  
25           embodiment, the degradation of the lipopeptide may be measured by HPLC, ultraviolet spectroscopy, infrared spectroscopy, NMR, or mass spectroscopy. In an even more preferred embodiment, HPLC is used to determine the percentage of intact lipopeptide that remains after a crystalline form of a lipopeptide has been subjected to a test of its stability.

30           Without wishing to be bound by any theory, applicants believe that daptomycin is crystallized by the methods described above. However, it is thought that washing and/or drying the daptomycin crystals causes the daptomycin crystalline material to revert to a non-crystalline but still crystal-like form. Nevertheless, even if the methods described

above only precipitate rather than crystallize the daptomycin or other lipopeptide, the methods still are advantageous because the methods purify the lipopeptide.

The invention also provides a crystalline or crystal-like lipopeptide produced by the above-described methods. In one embodiment, the crystalline or crystal-like lipopeptide comprises a lower amount of one or more impurities compared to the lipopeptide before crystallization or precipitation. In one embodiment, crystalline or crystal-like lipopeptide is daptomycin that comprises a lower level of anhydro-daptomycin and/or the  $\beta$ -isomer of daptomycin compared to daptomycin before crystallization or precipitation. In another embodiment, crystalline or crystal-like daptomycin comprises a lower level of all impurities compared to amorphous daptomycin. Similarly, in another embodiment, the crystalline or crystal-like lipopeptide is a daptomycin-related lipopeptide, as described above, which comprises a lower level of one or more impurities compared to an amorphous form of the daptomycin-related lipopeptide. In yet another embodiment, the crystalline or crystal-like daptomycin-related lipopeptide comprises a lower level of all impurities compared to an amorphous form of the daptomycin-related lipopeptide.

The crystalline or crystal-like lipopeptide produced by the method described above likely comprises monovalent or divalent cations and water. In a preferred embodiment, the crystalline or crystal-like lipopeptide is daptomycin or daptomycin-related lipopeptide that comprises a divalent cation. In a more preferred embodiment, the divalent cation is a calcium cation. In an even more preferred embodiment, the crystalline or crystal-like daptomycin or daptomycin-related lipopeptide comprises approximately 1-10% by weight of a divalent calcium cation and approximately 0-15% by weight of water as determined by atomic absorption or thermal gravity analysis. In a further preferred embodiment, the crystalline or crystal-like lipopeptide is daptomycin that comprises approximately 5% by weight of a divalent calcium cation and approximately 10% by weight of water; by HPLC analysis, the purity of the crystalline or crystal-like daptomycin is at least 95%, 96%, 97% or 98% or is any purity between 95-98%, relative to related substances and organic contaminants. Alternatively, the crystalline or crystal-like daptomycin or daptomycin-related lipopeptide comprises a monovalent cation such as sodium. Without wishing to be bound by any theory, it is thought that daptomycin or a daptomycin-related lipopeptide may form a salt with the monovalent or divalent cation when it crystallizes or precipitates.

The crystalline form of the lipopeptide may exhibit an increased solubility in a solution or an increased rate of reconstitution in a solution than an amorphous form of the lipopeptide. One may measure whether the crystalline lipopeptide exhibits an increased solubility or increased reconstitution rate by any method known in the art. For instance, one may dissolve a defined amount of a crystalline lipopeptide in an aqueous solution and measure the concentration of the dissolved lipopeptide and compare it to the concentration of dissolved lipopeptide that has been prepared by dissolving the same amount of amorphous lipopeptide in an aqueous solution. Similarly, one may measure the reconstitution rate of a crystalline lipopeptide by adding the crystalline lipopeptide to an aqueous solution and then measuring the concentration of dissolved lipopeptide over time and comparing it to the reconstitution rate of an amorphous lipopeptide that has been measured in the same way. The concentration of lipopeptide is measured by HPLC.

The methods described above provide for the production of crystalline or crystal-like lipopeptides that are more pure than the amorphous lipopeptide from which they are crystallized or precipitated. In one embodiment, the lipopeptide is daptomycin or a daptomycin-related lipopeptide. In another embodiment, daptomycin or a daptomycin-related lipopeptide has a purity of no more than 92% before crystallization and has a purity of at least approximately 95%, 96%, 97% or 98% purity, or any purity between 95-98%, after crystallization or precipitation as a crystal-like lipopeptide. In a still further preferred embodiment, daptomycin or a daptomycin-related lipopeptide has a purity of no more than 90% before crystallization and has a purity of approximately at least 97% or 98% after crystallization.

In another embodiment, the daptomycin has a purity of no more than 80%, preferably no more than 70% and more preferably no more than 60% purity before crystallization or precipitation, and has at least approximately 95%, 96%, 97% or 98% purity, or any purity between 95-98%, after purification. In another embodiment, the daptomycin has a purity of no more than 50%, preferably no more than 40%, more preferably no more than 30% purity before crystallization and has at least approximately 95%, 96%, 97% or 98% purity, or any purity between 95-98%, after purification by crystallization or precipitation. Further preferred is an embodiment in which daptomycin has a purity of no more than 20%, more preferably no more than 15%, even more preferably no more than 10% purity before crystallization and has at least approximately 95%, 96%, 97% or 98% purity, or any purity between 95-98%, after purification.

In a more preferred embodiment, the lipopeptide is daptomycin. A daptomycin preparation may be obtained by any method disclosed, e.g., in any one United States Patents RE32,333, RE32,455, 4,800,157, RE32,310, RE32,311, 4,537,717, 4,482,487, 4,524,135, 4,874,843, 4,885,243 or 5,912,226, which are herein incorporated specifically  
5 by reference. A daptomycin preparation may also be obtained by one of the methods described in International PCT Publication WO 01/53330, published July 26, 2001. After the lipopeptide preparation is prepared, the lipopeptide preparation is crystallized or precipitated following the teachings of the specification described herein to produce a crystalline or crystal-like lipopeptide that is more pure or that contains lower levels of  
10 specific impurities, e.g., anhydro-daptomycin, than the lipopeptide preparation from which it is prepared.

#### Processes for Producing Purified Lipopeptides from Fermentation Cultures

Another embodiment of the present invention is drawn to a process combining  
15 process chromatography steps and crystallization or precipitation to produce a purified lipopeptide. In a preferred embodiment, the method comprises the steps of producing a lipopeptide by any method known in the art, such as fermentation of a naturally-occurring or recombinant organism, and then subjecting the lipopeptide preparation to any one or more purification methods such as microfiltration, anion exchange chromatography,  
20 hydrophobic interaction chromatography, and/or size exclusion chromatography (either via traditional size exclusion chromatographic media or via ultrafiltration) to produce a lipopeptide preparation that has been partially purified, and then crystallizing or precipitating the lipopeptide preparation to obtain a purified crystalline or crystal-like lipopeptide. In a preferred embodiment, the lipopeptide is daptomycin or a daptomycin-  
25 related lipopeptide. The steps regarding fermentation, microfiltration, anion exchange chromatography, hydrophobic interaction chromatography and ultrafiltration are disclosed in the art, e.g., in any one United States Patents RE32,333, RE32,455, 4,800,157, RE32,310, RE32,311, 4,537,717, 4,482,487, 4,524,135, 4,874,843, 4,885,243 or 5,912,226, in International Publication WO 01/53330, published July 26, 2001.

30 The method optionally comprises the step of collecting and/or washing the crystalline or crystal-like material after the crystallization or precipitation step. In a preferred embodiment, the crystalline lipopeptide preparation may be collected by filtration. In another embodiment, the crystalline or crystal-like material is dried.

In one embodiment, the purification method comprises fermenting *Streptomyces roseosporus* to obtain a fermentation culture containing daptomycin. In one embodiment, the *S. roseosporus* may be fermented as described in United States Patent 4,885,243. In another embodiment, the fermentation conditions in which the A-21978C<sub>0</sub>-containing  
5 crude product is produced by *Streptomyces roseosporus* is altered in order to increase daptomycin production and decrease impurities and related contaminants produced by the *S. roseosporus* fermentation culture as described in International PCT Publication WO 01/53330, published July 26, 2001. The WO 01/53330 publication describes fermenting *S. roseosporus* as described in the '243 patent with the modification that the decanoic acid  
10 feed is kept at the lowest levels possible without diminishing the overall yield of the fermentation.

Alternatively, daptomycin may be obtained by fermenting a bacterial strain or other producing organism that recombinantly produces daptomycin. In one embodiment, the recombinant bacterial strain or other recombinant organism comprises the daptomycin  
15 biosynthetic gene cluster. In another embodiment, the daptomycin biosynthetic gene cluster or a portion thereof is introduced into the organism or bacterial strain via a bacterial artificial chromosome (BAC). In another embodiment, the recombinant bacterial strain used is *S. roseosporus* or *S. lividans* comprising a BAC containing the daptomycin biosynthetic gene cluster. United States Provisional Application 60/272,207,  
20 filed February 28, 2001 describes the daptomycin biosynthetic gene cluster from *S. roseosporus* and uses thereof, and is hereby incorporated by reference in its entirety.

After fermentation, the fermentation broth is clarified by centrifugation, microfiltration or extraction, as is known in the art or as described in the WO 01/53330 publication. In a preferred embodiment, the clarification is performed by microfiltration.  
25 See, e.g., Examples 13-16 and Figures 11-15. Figure 11 shows an exemplary manufacturing process that does not use crystallization or precipitation.

After the fermentation broth is clarified, the concentration of daptomycin in the broth is approximately 5-10%. In one embodiment of the invention, the daptomycin preparation is subjected to a crystallization/precipitation method described above directly  
30 subsequent to microfiltration. In one embodiment, crystallization or precipitation is performed under sterile conditions. After crystallization or precipitation is complete, the crystalline or crystal-like daptomycin is optionally collected, washed and dried, as



described in further detail below. The dry bulk active drug may then be used to dry fill sterile vials. See, e.g., Example 16 and Figure 12.

After clarification of the fermentation broth, the lipopeptide may be enriched in the preparation by anion exchange chromatography, as is known in the art or as described in the WO 01/53330 publication or herein. See, e.g., Examples 13-14 and Figures 12-13. After anion exchange chromatography, the purity of daptomycin in the broth is approximately 35-40%. In one embodiment of the invention, the daptomycin preparation is then subjected to a crystallization or precipitation method described above directly subsequent to anion exchange chromatography. In one embodiment, crystallization or precipitation is performed under sterile conditions. After crystallization or precipitation is complete, the crystalline or crystal-like daptomycin is optionally collected, washed and dried as described below. The dry bulk active drug may then be used to dry fill sterile vials. See, e.g., Example 14 and Figure 13.

In another embodiment of the invention, the daptomycin preparation is subjected to size exclusion ultrafiltration after anion exchange chromatography. Size exclusion ultrafiltration is described in the WO 01/53330 publication. The application published July 26, 2001 describes a method of depyrogenating, filtering and concentrating the daptomycin using an ultrafiltration membrane of 10,000 to 30,000 nominal molecular weight (NMW). The application discloses a method in which the lipopeptide passes through the ultrafiltration membrane while large molecular weight impurities, such as endotoxins, are retained by the filter. After the lipopeptide has passed through the membrane, the pH, temperature and/or salt concentration of the lipopeptide solution are altered such that the lipopeptides form micelles. The lipopeptide solution is then filtered on the ultrafiltration membrane under conditions in which the lipopeptide micelles are retained on the membrane while smaller impurities pass through the filter. In this manner, the lipopeptide is further purified. The application discloses the conditions under which lipopeptide micelles may be formed and disassociated as well as methods for filtering the lipopeptide solution to obtain a more purified lipopeptide application. In an even more preferred embodiment, the lipopeptide is daptomycin or a daptomycin-related lipopeptide. The lipopeptide may then be crystallized, as described herein. After both anion exchange chromatography and size exclusion ultrafiltration, daptomycin purity is approximately 80-90%. As discussed above, the daptomycin preparation is then subjected to a crystallization/precipitation method described above, preferably under

sterile conditions. The crystalline or crystal-like daptomycin may be optionally collected, washed, dried and used to dry fill vials as described below. See, e.g., Example 13 and Figure 12.

5 In another embodiment of the invention, the crude daptomycin preparation is subjected to size exclusion ultrafiltration without anion exchange chromatography. After size exclusion ultrafiltration, daptomycin purity is approximately 35-40%. The lipopeptide may then be crystallized or precipitated as described herein, preferably by sterile methods. As discussed above, the crystalline or crystal-like daptomycin may be collected, washed, dried and used to dry fill sterile vials. See, e.g., Example 15 and  
10 Figure 14.

In an alternative embodiment, the lipopeptide preparation is subjected to hydrophobic interaction chromatography (HIC), such as is described in the WO 01/53330 publication, after either the anion exchange chromatography or the size exclusion filtration. The lipopeptide may then be crystallized or precipitated as described herein.

15 After crystallization or precipitation, the crystalline or crystal-like lipopeptide may be collected by a method described herein, e.g., by filtration or centrifugation. The crystalline or crystal-like lipopeptide is optionally washed to remove residual crystallization or precipitation solvent. A method of washing crystals or crystal-like material are described below. See, e.g., Example 3. The washed or unwashed crystal or  
20 crystal-like material may be dried. The drying may be performed by any method known in the art, including, without limitation, vacuum drying, spray drying, tray drying or lyophilization. In one embodiment, the drying is performed under sterile conditions. In another embodiment, the drying is performed by vacuum drying. In a more preferred embodiment, the drying is performed using a 0.65 m<sup>3</sup> Klein Hastelloy-B double cone  
25 vacuum dryer or an equivalent apparatus. The dried crystalline or crystal-like lipopeptide is stable and is easily stored.

In one embodiment, vials are filled with any convenient amount of the dried crystalline or crystal-like lipopeptide. In one embodiment, the vials are filled under sterile conditions and then stoppered. In another embodiment, the vials are filled with 50  
30 to 5000 mg each of the dried crystalline or crystal-like lipopeptide. In another embodiment, the vials are filled with 100 to 1000 mg each. In another embodiment, the vials are filled with 200 to 500 mg each. In another embodiment, the dried crystalline or crystal-like lipopeptide is used for bulk packaging of the lipopeptide. The bulk packaging

is usually greater than 5000 mg each of the dried crystalline or crystal-like lipopeptide. In one embodiment, the bulk packaging is performed under sterile conditions.

In one embodiment, the crystallization or precipitation step is performed under sterile conditions. In this embodiment, sterile crystallization or precipitation reagents and a sterile, controlled working environment are used. In one embodiment, the lipopeptide is filtered on a ultrafiltration membrane, as disclosed above, before being mixed with the sterile crystallization/precipitation reagents. After crystallization or precipitation, the crystalline or crystal-like lipopeptide preparation is collected by centrifugation or filtration under sterile conditions. In one embodiment, the lipopeptide preparation is collected by sterile filtration. In another embodiment, the crystalline or crystal-like lipopeptide is sterilized after it has been collected. Methods of sterile crystallization, precipitation and filtration as well as methods of sterilizing a final pharmaceutical product are known in the art. See, e.g., Remington: The Science and Practice of Pharmacy, Easton, Pennsylvania: Mack Publishing Company (1995), pp. 1474-1487, herein incorporated by reference.

In another embodiment, the crystalline or crystal-like lipopeptide is not dried. In this embodiment, the crystalline or crystal-like lipopeptide is preferably stored in a solution that preserves the crystalline or crystal-like nature of the lipopeptide. Vials may be filled with the lipopeptide and solution under sterile or nonsterile conditions. In one embodiment, the conditions are sterile. Alternatively, the crystalline or crystal-like lipopeptide and solution may be used to fill bulk packaging.

Figures 10 and 11 provide flowcharts describing an exemplary daptomycin manufacturing protocol using crystallization. The incorporation of sterile crystallization into the manufacturing protocol shortens the protocol considerably and eliminates 3 to 4 steps in the process.

#### Crystalline or Crystal-like Lipopeptides, Pharmaceutical Compositions and Methods of Use Thereof

Another object of the instant invention is to provide crystalline or crystal-like lipopeptides or salts thereof, as well as pharmaceutical formulations comprising a crystalline or crystal-like lipopeptide or its salts. In one embodiment, the crystalline or crystal-like lipopeptide is daptomycin. However, all reference herein to crystalline or crystal-like lipopeptides specifically contemplates daptomycin, a daptomycin-related

molecule, including, *inter alia*, daptomycin, A54145 and a daptomycin-related lipopeptide, as disclosed above.

Daptomycin crystals or crystal-like particles, as well as other lipopeptide crystals or crystal-like particles may have a shape such as, *inter alia*, a needle-like shape, a plate-like shape, a lath-like shape, an equant-like shape, an urchin-like shape or a rod-like shape. In one embodiment, daptomycin crystals or crystal-like particles have an urchin-like, needle-like or rod-like shape. The size of the crystals or crystal-like particles may range from approximately 0.5  $\mu\text{m}$  to greater than 100  $\mu\text{m}$ . In one embodiment, the particle size is at least 5  $\mu\text{m}$  or greater. In a more preferred embodiment, the particle size is at least 10  $\mu\text{m}$  or greater, more preferably at least 50  $\mu\text{m}$ . In an even more preferred embodiment, the particle size is at least 100  $\mu\text{m}$ .

Further, in one embodiment, daptomycin crystals have an x-ray diffraction pattern as shown in Figs. 6, 7 and 8. In another embodiment, the lipopeptide crystal exhibits a different melting point than the amorphous form of the lipopeptide.

In one embodiment of the invention, a crystalline form of a lipopeptide exhibits a stability that is equal to or greater than the amorphous form of the lipopeptide. In a preferred embodiment, the crystalline form is daptomycin or a daptomycin-related lipopeptide. In another preferred embodiment, the crystalline lipopeptide is sterile. In another preferred embodiment, the stability of the crystalline lipopeptide is greater than the amorphous form of the lipopeptide. The crystalline lipopeptide may exhibit higher stability to heat, light, degradation or humidity than the amorphous form. The stability of the lipopeptide may be measured by any means including, e.g., antibiotic activity, degradation of the lipopeptide or conversion of daptomycin to anhydro-daptomycin or the  $\beta$ -isomer of daptomycin. In another embodiment of the invention, the crystalline form of the lipopeptide may be more quickly reconstituted in aqueous solution than the amorphous form of the lipopeptide.

Crystalline or crystal-like lipopeptides, such as daptomycin or a daptomycin-related lipopeptide, pharmaceutically-acceptable salts, esters, amides, ethers and protected forms thereof, can be formulated for oral, intravenous, intramuscular, subcutaneous, aerosol, topical or parenteral administration for the therapeutic, empirical or prophylactic treatment of diseases, particularly bacterial infections. Reference herein to "crystalline or crystal-like lipopeptides" or "crystalline or crystal-like daptomycin" includes

pharmaceutically acceptable salts thereof. Crystalline or crystal-like lipopeptides, such as daptomycin, may be particularly advantageous for pharmaceutical compositions because they can be easily formulated as micronized particles of microspheres, which permits the facile preparation of enterically coated lipopeptides for oral delivery, pharmaceutical compositions for aerosol delivery to, e.g., the lung, and the preparation of lipopeptides formulations for sustained release. Crystalline or crystal-like lipopeptides and crystalline or crystal-like daptomycin may also be more readily dissolved in aqueous solution.

Crystalline or crystal-like lipopeptides, including daptomycin or daptomycin-related lipopeptides can be formulated using any pharmaceutically acceptable carrier or excipient that is compatible with daptomycin or with the lipopeptide of interest. See, e.g., Handbook of Pharmaceutical Additives: An International Guide to More than 6000 Products by Trade Name, Chemical, Function, and Manufacturer, Ashgate Publishing Co., eds., M. Ash and I. Ash, 1996; The Merck Index: An Encyclopedia of Chemicals, Drugs and Biologicals, ed. S. Budavari, annual; Remington's Pharmaceutical Sciences, Mack Publishing Company, Easton, PA; Martindale: The Complete Drug Reference, ed. K. Parfitt, 1999; and Goodman & Gilman's The Pharmaceutical Basis of Therapeutics, Pergamon Press, New York, NY, ed. L. S. Goodman et al.; the contents of which are incorporated herein by reference, for a general description of the methods for administering various antimicrobial agents for human therapy. Compounds of this invention can be mixed with conventional pharmaceutical carriers and excipients and used in the form of tablets, capsules, elixirs, suspensions, syrups, wafers, creams and the like. Compounds of this invention may also be mixed with other therapeutic agents and antibiotics, such as discussed herein. The compositions comprising a compound of this invention will contain from about 0.1 to about 90% by weight of the active compound, and more generally from about 10 to about 30%.

The compositions of the invention can be delivered using controlled (e.g., capsules) or sustained release delivery systems (e.g., bioerodable matrices). Exemplary delayed release delivery systems for drug delivery that are suitable for administration of the compositions of the invention are described in U.S. Patent Nos. 4,452,775 (issued to Kent), 5,239,660 (issued to Leonard), 3,854,480 (issued to Zaffaroni).

The compositions may contain common carriers and excipients, such as corn starch or gelatin, lactose, sucrose, microcrystalline cellulose, kaolin, mannitol, dicalcium phosphate, sodium chloride and alginic acid. The compositions may contain

croscarmellose sodium, microcrystalline cellulose, corn starch, sodium starch glycolate and alginic acid.

Tablet binders that can be included are acacia, methylcellulose, sodium carboxymethylcellulose, polyvinylpyrrolidone (Povidone), hydroxypropyl methylcellulose, sucrose, starch and ethylcellulose.

Lubricants that can be used include magnesium stearate or other metallic stearates, stearic acid, silicone fluid, talc, waxes, oils and colloidal silica.

Flavoring agents such as peppermint, oil of wintergreen, cherry flavoring or the like can also be used. It may also be desirable to add a coloring agent to make the dosage form more aesthetic in appearance or to help identify the product.

For oral use, solid formulations such as tablets and capsules are particularly useful. Sustained release or enterically coated preparations may also be devised. In another embodiment, crystalline or crystal-like lipopeptides may be supplied in combination with a carrier composition that enhances the oral availability of the lipopeptide. In a preferred embodiment, the crystalline or crystal-like lipopeptide is daptomycin. For pediatric and geriatric applications, suspensions, syrups and chewable tablets are especially suitable. For oral administration, the pharmaceutical compositions are in the form of, for example, a tablet, capsule, suspension or liquid. The pharmaceutical composition is preferably made in the form of a dosage unit containing a therapeutically-effective amount of the active ingredient. Examples of such dosage units are tablets and capsules. For therapeutic purposes, the tablets and capsules which can contain, in addition to the active ingredient, conventional carriers such as binding agents, for example, acacia gum, gelatin, polyvinylpyrrolidone, sorbitol, or tragacanth; fillers, for example, calcium phosphate, glycine, lactose, maize-starch, sorbitol, or sucrose; lubricants, for example, magnesium stearate, polyethylene glycol, silica, or talc; disintegrants, for example, potato starch, flavoring or coloring agents, or acceptable wetting agents. Oral liquid preparations generally are in the form of aqueous or oily solutions, suspensions, emulsions, syrups or elixirs may contain conventional additives such as suspending agents, emulsifying agents, non-aqueous agents, preservatives, coloring agents and flavoring agents. Oral liquid preparations may comprise lipopeptide micelles or monomeric forms of the lipopeptide. Examples of additives for liquid preparations include acacia, almond oil, ethyl alcohol, fractionated coconut oil, gelatin,

glucose syrup, glycerin, hydrogenated edible fats, lecithin, methyl cellulose, methyl or propyl *para*-hydroxybenzoate, propylene glycol, sorbitol, or sorbic acid.

For intravenous (IV) use, a water soluble form of a compound of this invention can be dissolved in any of the commonly used intravenous fluids and administered by  
5 infusion. Intravenous formulations may include carriers, excipients or stabilizers including, without limitation, calcium, human serum albumin, citrate, acetate, calcium chloride, carbonate, and other salts. Intravenous fluids include, without limitation, physiological saline or Ringer's solution. Daptomycin or other lipopeptides also may be placed in injectors, cannulae, catheters and lines.

10 Formulations for parenteral administration can be in the form of aqueous or non-aqueous isotonic sterile injection solutions or suspensions. These solutions or suspensions can be prepared from sterile powders or granules having one or more of the carriers mentioned for use in the formulations for oral administration. The crystalline or crystal-like lipopeptides can be dissolved in polyethylene glycol, propylene glycol,  
15 ethanol, corn oil, benzyl alcohol, sodium chloride, and/or various buffers. For intramuscular, parenteral or intravenous preparations, a sterile formulation of a crystalline or crystal-like lipopeptide compound or a suitable soluble salt form of the compound, for example the hydrochloride salt, can be dissolved and administered in a pharmaceutical diluent such as Water-for-Injection (WFI), physiological saline or 5% glucose. A suitable  
20 insoluble form of the crystalline or crystal-like lipopeptide also may be prepared and administered as a suspension in an aqueous base or a pharmaceutically acceptable oil base, e.g., an ester of a long chain fatty acid such as ethyl oleate.

Injectable depot forms may be made by forming microencapsulated matrices of the crystalline or crystal-like lipopeptide in biodegradable polymers such as polylactide-  
25 polyglycolide. Depending upon the ratio of drug to polymer and the nature of the particular polymer employed, the rate of drug release can be controlled. Examples of other biodegradable polymers include poly(orthoesters) and poly(anhydrides). Depot injectable formulations are also prepared by entrapping the drug in microemulsions that are compatible with body tissues.

30 For topical use the compounds of the present invention can also be prepared in suitable forms to be applied to the skin, or mucus membranes of the nose and throat, and can take the form of creams, ointments, liquid sprays or inhalants, lozenges, or throat paints. Such topical formulations further can include chemical compounds such as

dimethylsulfoxide (DMSO) to facilitate surface penetration of the active ingredient. For topical preparations, a sterile formulation comprising a crystalline or crystal-like lipopeptide, such as crystalline or crystal-like daptomycin, a suitable salt form thereof, may be administered in a cream, ointment, spray or other topical dressing. Topical  
5 preparations may also be in the form of bandages that have been impregnated with a lipopeptide composition.

For application to the eyes or ears, the compounds of the present invention can be presented in liquid or semi-liquid form formulated in hydrophobic or hydrophilic bases as ointments, creams, lotions, paints or powders.

10 For rectal administration the compounds of the present invention can be administered in the form of suppositories admixed with conventional carriers such as cocoa butter, wax or other glyceride.

For aerosol preparations, a sterile formulation of a crystalline or crystal-like lipopeptide or a salt form of the compound may be used in inhalers, such as metered dose  
15 inhalers, and nebulizers. Aerosolized forms may be especially useful for treating respiratory infections, such as pneumonia and sinus-based infections.

Alternatively, the compounds of the present invention can be in powder crystalline or crystal-like form for reconstitution in the appropriate pharmaceutically acceptable carrier at the time of delivery. In another embodiment, the unit dosage form of the  
20 compound can be a solution of the compound or a salt thereof in a suitable diluent in sterile, hermetically sealed ampules. The concentration of the compound in the unit dosage may vary, e.g. from about 1 percent to about 50 percent, depending on the compound used and its solubility and the dose desired by the physician. If the compositions contain dosage units, each dosage unit preferably contains approximately  
25 from 10-5000 mg of the active material, more preferably 50 to 1000 mg, and even more preferably 100 to 500 mg. For adult human treatment, the dosage employed preferably ranges from 100 mg to 3 g, per day, depending on the route and frequency of administration.

In a further aspect, this invention provides a method for treating an infection  
30 caused by a gram-positive bacteria in a subject. In a preferred embodiment, the method may be used to treat an infection caused by a gram-positive bacteria. The term "treating" is defined as administering, to a subject, a therapeutically-effective amount of a compound of the invention, both to prevent the occurrence of an infection and to control



or eliminate an infection, e.g., an established infection. The term "subject", as described herein, is defined as a mammal, a plant or a cell culture. As used herein, the phrase "therapeutically-effective amount" means an amount of daptomycin, daptomycin-related lipopeptide or other antibacterial lipopeptide according to the present invention that prevents the onset, alleviates the symptoms, or stops the progression of a bacterial infection. In a preferred embodiment, a subject is a human or other animal patient in need of lipopeptide treatment. An established infection may be one that is acute or chronic. An effective dose is generally between about 0.1 and about 75 mg/kg crystalline or crystal-like lipopeptide, such as crystalline or crystal-like daptomycin or daptomycin-related lipopeptide, or a pharmaceutically acceptable salt thereof. A preferred dose is from about 1 to about 25 mg/kg of crystalline or crystal-like daptomycin or daptomycin-related lipopeptide or a pharmaceutically acceptable salt thereof. A more preferred dose is from about 1 to 12 mg/kg crystalline or crystal-like daptomycin, a crystalline or crystal-like daptomycin-related lipopeptide or a pharmaceutically acceptable salt thereof. An even more preferred dose is about 3 to 8 mg/kg crystalline or crystal-like daptomycin or daptomycin-related lipopeptide or a pharmaceutically acceptable salt thereof. Exemplary procedures for delivering an antibacterial agent are described in U.S. Patent No. 5,041,567, issued to Rogers and in International PCT Publication WO 95/05384, the entire contents of which documents are incorporated in their entirety herein by reference.

The crystalline or crystal-like lipopeptide, e.g., daptomycin, can be administered as a single daily dose or in multiple doses per day. The treatment regime may require administration over extended periods of time, e.g., for several days or for from two to four weeks. The amount per administered dose or the total amount administered will depend on such factors as the nature and severity of the infection, the age and general health of the patient, the tolerance of the patient to the lipopeptide and the microorganism or microorganisms involved in the infection. A method of administration is disclosed in WO 00/18419, published April 6, 2000, herein incorporated by reference.

The methods of the present invention comprise administering a compound of the invention, or a pharmaceutical composition thereof to a patient in need thereof in an amount that is efficacious in reducing or eliminating the gram-positive bacterial infection. The lipopeptide may be administered orally, parenterally, by inhalation, topically, rectally, nasally, buccally, vaginally, or by an implanted reservoir, external pump or catheter. The lipopeptide may be prepared for ophthalmic or aerosolized uses.

Compounds of the invention, or pharmaceutical compositions thereof also may be directly injected or administered into an abscess, ventricle or joint. Parenteral administration includes subcutaneous, intravenous, intramuscular, intra-articular, intra-synovial, cisternal, intrathecal, intrahepatic, intralesional and intracranial injection or infusion. In a preferred embodiment, crystalline or crystal-like daptomycin, daptomycin-related lipopeptide or other lipopeptide is administered intravenously, subcutaneously or orally.

The method of the instant invention may be used to treat a patient having a bacterial infection in which the infection is caused or exacerbated by any type of gram-positive bacteria. In a preferred embodiment, crystalline or crystal-like daptomycin, daptomycin-related lipopeptide or other lipopeptide, or pharmaceutical compositions thereof, are administered to a patient according to the methods of this invention. In another embodiment, the bacterial infection may be caused or exacerbated by bacteria including, but not limited to, methicillin-susceptible and methicillin-resistant staphylococci (including *Staphylococcus aureus*, *Staphylococcus epidermidis*, *Staphylococcus haemolyticus*, *Staphylococcus hominis*, *Staphylococcus saprophyticus*, and coagulase-negative staphylococci), glycopeptide intermediary- susceptible *Staphylococcus aureus* (GISA), penicillin-susceptible and penicillin-resistant streptococci (including *Streptococcus pneumoniae*, *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus avium*, *Streptococcus bovis*, *Streptococcus lactis*, *Streptococcus sanguis* and *Streptococci* Group C, *Streptococci* Group G and viridans streptococci), enterococci (including vancomycin-susceptible and vancomycin-resistant strains such as *Enterococcus faecalis* and *Enterococcus faecium*), *Clostridium difficile*, *Clostridium clostridiiforme*, *Clostridium innocuum*, *Clostridium perfringens*, *Clostridium ramosum*, *Haemophilus influenzae*, *Listeria monocytogenes*, *Corynebacterium jeikeium*, *Bifidobacterium* spp., *Eubacterium aerofaciens*, *Eubacterium lentum*, *Lactobacillus acidophilus*, *Lactobacillus casei*, *Lactobacillus plantarum*, *Lactococcus* spp., *Leuconostoc* spp., *Pediococcus*, *Peptostreptococcus anaerobius*, *Peptostreptococcus asaccarolyticus*, *Peptostreptococcus magnus*, *Peptostreptococcus micros*, *Peptostreptococcus prevotii*, *Peptostreptococcus productus*, *Propionibacterium acnes*, and *Actinomyces* spp.

The antibacterial activity of daptomycin against classically "resistant" strains is comparable to that against classically "susceptible" strains in *in vitro* experiments. In addition, the minimum inhibitory concentration (MIC) value for daptomycin against

susceptible strains is typically 4-fold lower than that of vancomycin. Thus, in a preferred embodiment, a compound of the invention, or a pharmaceutical composition of any one of these crystalline or crystal-like lipopeptides, is administered according to the methods of this invention to a patient who exhibits a bacterial infection that is resistant to other antibiotics, including vancomycin. In addition, unlike glycopeptide antibiotics, daptomycin exhibits rapid, concentration-dependent bactericidal activity against gram-positive organisms. Thus, in a preferred embodiment, compounds of the invention, or a pharmaceutical composition of any one of these crystalline or crystal-like lipopeptides, is administered according to the methods of this invention to a patient in need of rapidly acting antibiotic therapy.

The method of the instant invention may be used for a gram-positive bacterial infection of any organ or tissue in the body. These organs or tissue include, without limitation, skeletal muscle, skin, bloodstream, kidneys, heart, lung and bone. The method of the invention may be used to treat, without limitation, skin and soft tissue infections, bacteremia and urinary tract infections. The method of the invention may be used to treat community acquired respiratory infections, including, without limitation, otitis media, sinusitis, chronic bronchitis and pneumonia, including pneumonia caused by drug-resistant *Streptococcus pneumoniae* or *Haemophilus influenzae*. The method of the invention also may be used to treat mixed infections that comprise different types of gram-positive bacteria, including aerobic, caprophilic or anaerobic bacteria. These types of infections include intra-abdominal infections, pneumonia, bone and joint infections and obstetrical/gynecological infections. The method of the invention also may be used to treat an infection including, without limitation, endocarditis, nephritis, septic arthritis and osteomyelitis. In a preferred embodiment, any of the above-described diseases may be treated using crystalline or crystal-like daptomycin, daptomycin-related lipopeptide, antibacterial lipopeptide, or pharmaceutical compositions of any one of these crystalline or crystal-like lipopeptides.

Crystalline or crystal-like daptomycin, daptomycin-related lipopeptide or other lipopeptide may also be administered in the diet or feed of a patient or animal. If administered as part of a total dietary intake, the amount of daptomycin or other lipopeptide can be less than 1% by weight of the diet and preferably no more than 0.5% by weight. The diet for animals can be normal foodstuffs to which daptomycin or other lipopeptide can be added or it can be added to a premix.

The method of the instant invention may also be practiced while concurrently administering another form of daptomycin or other lipopeptide antibiotic, e.g., one that is not crystalline or crystal-like, or with one or more antifungal agents and/or one or more antibiotics other than crystalline or crystal-like daptomycin or other crystalline or crystal-like lipopeptide antibiotics. Co-administration of an antifungal agent and an antibiotic other than crystalline or crystal-like daptomycin or another lipopeptide antibiotic may be useful for mixed infections such as those caused by different types of gram-positive bacteria, or those that caused by both bacteria and fungus. Furthermore, crystalline or crystal-like daptomycin or other lipopeptide antibiotic may improve the toxicity profile of one or more co-administered antibiotics. It has been shown that administration of daptomycin and an aminoglycoside may ameliorate renal toxicity caused by the aminoglycoside. In a preferred embodiment, an antibiotic and/or antifungal agent may be administered concurrently with a compound of this invention, or in a pharmaceutical composition comprising a compound of this invention.

Antibacterial agents and classes thereof that may be co administered with a compound of the present invention include, without limitation, penicillins and related drugs, carbapenems, cephalosporins and related drugs, aminoglycosides, bacitracin, gramicidin, mupirocin, chloramphenicol, thiamphenicol, fusidate sodium, lincomycin, clindamycin, macrolides, novobiocin, polymyxins, rifamycins, spectinomycin, tetracyclines, vancomycin, teicoplanin, streptogramins, anti-folate agents including sulfonamides, trimethoprim and its combinations and pyrimethamine, synthetic antibacterials including nitrofurans, methenamine mandelate and methenamine hippurate, nitroimidazoles, quinolones, fluoroquinolones, isoniazid, ethambutol, pyrazinamide, para-aminosalicylic acid (PAS), cycloserine, capreomycin, ethionamide, prothionamide, thiacetazone, viomycin, everninomycin, glycopeptide, glycyclcycline, ketolides, oxazolidinone; imipenen, amikacin, netilmicin, fosfomycin, gentamicin, ceftriaxone, Zircin, LY 333328, CL 331002, HMR 3647, Linezolid, Synercid, Aztreonam, and Metronidazole, Epiroprim, OCA\_983, GV\_143253, Sanfetrinem sodium, CS\_834, Biapenem, A\_99058.1, A\_165600, A\_179796, KA 159, Dynemicin A, DX8739, DU 6681; Cefluprenam, ER 35786, Cefoselis, Sanfetrinem celexetil, HGP\_31, Cefpirome, HMR\_3647, RU\_59863, Mersacidin, KP 736, Rifalazil; Kosan, AM 1732, MEN 10700, Lenapenem, BO 2502A, NE\_1530, PR 39, K130, OPC 20000, OPC 2045, Venepriam, PD 138312, PD 140248, CP 111905, Sulopenem, ritipenam acoxyl, RO\_65\_5788,

Cyclotrialdine, Sch\_40832, SEP\_132613, micacocidin A, SB\_275833, SR\_15402, SUN A0026, TOC 39, carumonam, Cefozopran, Cefetamet pivoxil, and T 3811.

In a preferred embodiment, antibacterial agents that may be co administered with a compound according to this invention include, without limitation, imipenen, amikacin, 5 netilmicin, fosfomycin, gentamicin, ceftriaxone, teicoplanin, Ziracin, LY 333328, CL 331002, HMR 3647, Linezolid, Synercid, Aztreonam, and Metronidazole.

Antifungal agents that may be co administered with a compound according to this invention include, without limitation, Caspofungen, Voriconazole, Sertaconazole, IB\_367, FK\_463, LY\_303366, Sch\_56592, Sitafloxacin, DB\_289 polyenes, such as 10 Amphotericin, Nystatin, Primaricin; azoles, such as Fluconazole, Itraconazole, and Ketoconazole; allylamines, such as Naftifine and Terbinafine; and anti-metabolites such as Flucytosine. Other antifungal agents include without limitation, those disclosed in Fostel et al., Drug Discovery Today 5:25\_32 (2000), herein incorporated by reference. Fostel et al. disclose antifungal compounds including Corynecandin, Mer\_WF3010, 15 Fusacandins, Artrichitin/LL 15G256(, Sordarins, Cispentacin, Azoxybacillin, Aureobasidin and Khafrefungin.

Compounds of this invention, or a pharmaceutical composition of any one or more of these crystalline or crystal-like lipopeptides, may be administered according to this method until the bacterial infection is eradicated or reduced. In one embodiment, the 20 crystalline or crystal-like lipopeptide is administered for a period of time from approximately 3 days to approximately 6 months. In a preferred embodiment, the crystalline or crystal-like lipopeptide is administered for 7 to 56 days. In a more preferred embodiment, the crystalline or crystal-like lipopeptide is administered for 7 to 28 days. In an even more preferred embodiment, the crystalline or crystal-like lipopeptide is 25 administered for 7 to 14 days. The crystalline or crystal-like lipopeptide may be administered for a longer or shorter time period if it is so desired. In a preferred embodiment, the lipopeptide is daptomycin or daptomycin-related lipopeptide.

In order that this invention may be more fully understood, the following examples are set forth. These examples are for the purpose of illustration only and are not to be 30 construed as limiting the scope of the invention in any way.

#### EXAMPLE 1

Daptomycin was prepared by conventional techniques. The daptomycin preparation was a pale yellow amorphous powder, with a solubility at 25°C of greater than 1 g/mL in water and a solubility of 2.8 mg/mL in ethanol. The amorphous daptomycin preparation was hygroscopic and decomposed at 215°C.

5        The remaining examples describe crystallizing or precipitating lipopeptides in the presence or absence of an organic precipitant (e.g., PEG).

#### EXAMPLE 2

10        In a microbatch crystallization, 25 µL of a daptomycin stock (20 mg/mL in methanol) was sequentially mixed with 15 µL of reagent stock (200 mM calcium acetate, 0.1 M cacodylate (pH 6.5), 18% [w/v] PEG 8000 and 15 µL ethylene glycol) to give a solution that was 27.5% aqueous component, 45% methanol and 27.5% ethylene glycol. Urchin-like crystals were formed at a yield of 50% with a purity of 98% as measured by HPLC.

15

#### EXAMPLE 3

20        A daptomycin stock was prepared by dissolving 440 mg daptomycin in 1 mL of a buffer containing 25 mM sodium acetate (pH 5.0) and 5 mM CaCl<sub>2</sub>. Crystallization was done by the vapor diffusion (hanging drop) method, in which 5 µL of the daptomycin stock was added to 5 µL of 0.1 M tri-sodium citrate dihydrate (pH 5.6), and 35% [v/v] tert-butanol in water to form a drop. The drop was suspended over a reservoir solution (0.1 M tri-sodium citrate dihydrate (pH 5.6), and 35% [v/v] tert-butanol in water) in an air-tight environment until crystallization occurred. This method yielded urchin-like daptomycin crystals. See, e.g., Fig. 2.

25

#### EXAMPLE 4

30        5 µL of a daptomycin stock prepared as in Example 3 was added to 5 µL of a solution containing 0.1 M sodium cacodylate (pH 6.5), 0.2 M calcium acetate and 9% [w/v] PEG 8000. Crystallization was done by the vapor diffusion method as described in Example 3. This method yielded needle-like daptomycin crystals. See, e.g., Fig. 3.

#### EXAMPLE 5

5  $\mu$ L of a daptomycin stock prepared as in Example 3 was added to 5  $\mu$ L of a solution of 0.1 M sodium cacodylate (pH 6.5), 0.2 M zinc acetate and 9% [w/v] PEG 8000 containing 0.1  $\mu$ L benzamidine to give a final concentration of 220 mg/mL daptomycin. Crystallization was done by the vapor diffusion method as described in Example 3. This method yielded rod-like daptomycin crystals. See, e.g., Fig. 4.

#### EXAMPLE 6

One mL of daptomycin (97.1% pure as determined by HPLC) at a concentration of 20-25 mg/mL in water was sequentially mixed with 231  $\mu$ L water, 77  $\mu$ L of calcium acetate (pH 6.0), 960  $\mu$ L propylene glycol and 231  $\mu$ L of 50% [w/v] PEG 4000. The solution was allowed to sit for 4-5 hours at 4°C. Urchin-like crystals were formed at a yield of 75%. The crystalline daptomycin was washed with isopropanol. The daptomycin was 98.4% pure as determined by HPLC.

#### EXAMPLE 7

Daptomycin (200 mg, 97.1% pure) was dissolved in 2.54 mL water. The daptomycin solution was sequentially mixed in order with 10.0 mL methanol, 0.78 mL 1 M calcium acetate (pH 6.0), 9.50 mL propylene glycol and 2.20 mL 50% [w/v] PEG 4000 to give a final volume of 25.02 mL. The mixture was tumbled at room temperature for 10-14 hours in a hematology mixer (Fischer). Crystals began to appear within a few hours. Final yield was approximately 70-80% after 14 hours. The crystals were harvested by centrifugation at 1000 rpm for 15 minutes. The supernatant was removed and the crystals were resuspended in 12.5 mL isopropanol. The daptomycin suspension was transferred to a column (Biorad) and the isopropanol was removed by allowing it to drip by gravity. The crystals were dried by a nitrogen stream. Any lumps were broken up during the drying procedure to obtain a uniform dry sample. Crystals prepared by this method were urchin-like and had a purity of 98.37%.

#### EXAMPLE 8

Daptomycin was crystallized according to Example 7 except that PEG 8000 was used in replacement of PEG 4000. The quantities of reagents used are identical to those

in Example 7. Crystals prepared by this method were urchin-like and had a purity of 98.84%.

#### EXAMPLE 9

5 Two daptomycin samples prepared according to Example 7 and one amorphous sample were analyzed for crystallinity using the USP <695> crystallinity test. Daptomycin particles were mounted in mineral oil on a glass slide and then were examined by polarizing light microscope (PLM). The particles were determined to be crystalline if they were birefringent (have interference colors) and had extinction  
10 positions when the stage was rotated.

The amorphous daptomycin sample consisted of lacy, flaky particles that were not birefringent. There were a few sliver-like areas in some of the flakes that had weak birefringence, but the particles were primarily amorphous. In contrast, the daptomycin samples prepared according to Example 7 consisted of polycrystalline particles with weak  
15 birefringence and some extinction, indicating that they were primarily crystalline. See Fig. 5.

#### EXAMPLE 10

20 Two daptomycin samples prepared according to Example 7 and one amorphous sample were analyzed for crystallinity by x-ray powder diffraction. The samples were analyzed on a Siemens D500 Automated Powder Diffractometer (ORS 1D No. LD-301-4), which was operated according to ORS Standard Operation Procedure EQ-27 Rev. 9. The diffractometer was equipped with a graphite monochromator and a Cu ( $\lambda=1.54 \text{ \AA}$ ) x-ray source operated at 50 kV, 40 mA. Two-theta ( $\theta$ ) calibration is performed using an  
25 NBS mica standard (SRM675). The samples were analyzed using the following instrument parameters:

	Measuring Range for $2\theta$ (degrees)	4.0 - 40.0
	Step Width (degrees)	0.05
	Measuring Time per Step (secs)	1.2
30	Beam Slits	1(1°), 2(1°), 3(1°), 4(0.15°), 5(0.15°).

Sample preparation was performed according to ORS Standard Operation Procedure MIC-7 Rev. 1 using a zero background sample plate.



All samples were done using a Cu ( $\lambda=1.54 \text{ \AA}$ ) x-ray source. The amorphous daptomycin sample did not show any peaks by x-ray powder diffraction. See Fig. 6. In contrast, the two daptomycin samples both showed peaks by x-ray powder diffraction. The diffraction angle ( $2\theta$ ) of the first daptomycin sample (Fig. 7) was 19.225, 23.242, 23.427 and 23.603 (degree). The diffraction angle ( $2\theta$ ) for the second daptomycin sample (Fig. 8) was 10.966, 19.205 and 23.344 (degree). The first crystalline daptomycin sample also showed a small peak between 10-11°. See Fig. 7.

#### EXAMPLE 11

Daptomycin was dissolved in water. Sodium acetate was added to achieve a final concentration of 187 mM. Calcium chloride was added to achieve a final concentration of 28 mM. The daptomycin solution was mixed and isopropanol was added to a final concentration of 78.4%. The solution was mixed and incubated. A precipitated material was formed after incubation. The precipitated material appeared to be urchin-like crystals of approximately 60  $\mu\text{m}$  diameter by optical microscopy. The material was then dried. The dry material contained approximately 30-40% salt. After drying, powder x-ray diffraction was performed. The powder x-ray diffraction did not show the presence of crystals in the dried daptomycin precipitate.

#### EXAMPLE 12

One gram of daptomycin (approximately 91.5% purity as measured by HPLC) was added to 16.8 mL of distilled water and dissolved. 2.5 mL of 1M calcium acetate (pH 6.1) and 60 mL of isopropanol was added. The solution was placed in a 27°C water bath and permitted to equilibrate to temperature of the water bath. 5 mL aliquots of isopropanol were slowly added until the solution became cloudy (a total of approximately 30 mL isopropanol). The solution was incubated overnight at 27°C to form a precipitate. The precipitate appeared to contain urchin-like crystals of approximately 60  $\mu\text{m}$  by optical microscopy. See Figure 2.

The daptomycin precipitate was poured into a pressure filter/drying funnel and filtered by gravity. The precipitate was washed twice with 25 mL each time of a washing solution (80% isopropanol and 20% solution A where solution A consists of 18mL of water and 2mL of glacial acetic acid) and allowed to drip by gravity overnight. The

precipitate was then transferred to a desiccator and dried under vacuum. After drying, powder x-ray diffraction was performed. The powder x-ray diffraction did not show the presence of crystals in the dried daptomycin precipitate. However, purity analysis of the precipitated material by HPLC showed that the material was 98.2% pure daptomycin.

5 Significantly, the daptomycin preparation after precipitation has significantly less anhydro-daptomycin than the daptomycin preparation before precipitation.

Without wishing to be bound by any theory, applicants believe that the conditions used to precipitate the daptomycin in Examples 11 and 12 actually produce a crystalline form of daptomycin but that the subsequent washing steps and/or drying steps cause the  
10 crystalline daptomycin to revert to a non-crystalline form. Nonetheless, the non-crystalline daptomycin is still crystal-like as shown in Figure 3 by the birefringence of a crystal sample in polarized light.

### EXAMPLE 13

15 A fermentation culture of *S. roseosporus* NRRL Strain 15998 is conducted in a controlled decanoic acid feed fermentation at levels that optimize the production of the antibiotic while minimizing the production of contaminants. The residual decanoic acid feed is measured by gas chromatography and the target residual level is 10 ppm decanoic acid from the start of induction (approximately at hour 30) until harvest. Centrifugation  
20 of the culture and subsequent analysis of the clarified broth are used to measure the production of daptomycin by HPLC. The harvest titer is typically between 1.0 and 3.0 grams per liter of fermentation broth.

The fermentation culture is harvested either by microfiltration using a Pall-Sep or equivalent microfiltration system, or by full commercial-scale centrifugation and depth  
25 filter. The clarified broth is applied to an anion exchange resin, Mitsubishi FP-DA 13, washed with of 30 mM NaCl at pH 6.5 and eluted with of 300 mM NaCl at pH 6.0-6.5. Alternatively, the FP-DA 13 column is washed with of 30 mM NaCl at pH 6.5 and eluted with of 300 mM NaCl at pH 6.0-6.5. The pH is adjusted to 3.0-4.8 and the temperature is adjusted to 2-15°C. Under these conditions, daptomycin forms a micelle. The  
30 micellar daptomycin solution is filtered-washed using a 10,000 NMW ultrafilter (AG Technology Corp. UF hollow fiber or equivalent) in any configuration. The daptomycin micelles are retained by the filter, but a large number of impurities are eliminated because

they pass through the 10,000 NMW filter. Ultrafiltration of daptomycin micelles increases daptomycin purity to approximately 80-90%.

The daptomycin preparation is then crystallized or precipitated under sterile conditions using one of the methods described above. In a preferred embodiment, the daptomycin is crystallized or precipitated according to the protocol described in Examples 7, 8 or 12 except that it can be scaled up for large preparation of daptomycin. The crystalline or crystal-like daptomycin is separated from the crystallization/precipitation solution by filtration, preferably by vacuum filtration. The crystalline or crystal-like daptomycin is washed with washing solution (see Example 3). The crystalline or crystal-like daptomycin is then vacuum dried under sterile conditions using a 0.65 m<sup>3</sup> Klein Hastelloy-B double cone vacuum dryer or equivalent apparatus. Vials are then filled with either 250 or 500 mg of dried crystalline daptomycin per vial. Figure 9 shows a flowchart of this manufacturing method.

#### EXAMPLE 14

Fermentation of *S. roseosporus*, microfiltration of the fermentation culture and anion exchange chromatography is performed as described in Example 13. The daptomycin preparation is approximately 35-40% pure at this point. After anion exchange chromatography, the daptomycin is crystallized or precipitated according to the protocol described in Example 13. The daptomycin is then washed and dried according to the protocol set forth in Example 13. The dried crystalline or crystal-like daptomycin is then used to fill sterile vials as described in Example 13. Figure 6 shows a flowchart of this manufacturing method.

#### EXAMPLE 15

Fermentation of *S. roseosporus* and microfiltration of the fermentation culture is performed as described in Example 13. After microfiltration, the fermentation culture is subjected to size exclusion ultrafiltration as described in Example 13. The daptomycin preparation is approximately 35-40% pure at this point. After ultrafiltration, the daptomycin is crystallized or precipitated according to the protocol described in Example 13. The daptomycin is then washed and dried according to the protocol set forth in Example 13. The dried crystalline or crystal-like daptomycin is then used to fill sterile

vials as described in Example 13. Figure 7 shows a flowchart of this manufacturing method.

#### EXAMPLE 16

5 Fermentation of *S. roseosporus* and microfiltration of the fermentation culture is performed as described in Example 13. The daptomycin preparation is 5-10% pure at this point. After microfiltration, the fermentation culture is crystallized or precipitated according to the protocol described in Example 13. The daptomycin is then washed and dried and used to fill sterile vials as described in Example 13. Figure 8 shows a flowchart  
10 of this manufacturing method.

#### EXAMPLE 17

CB-131547 (see Figure x), a cyclic lipopeptide analog of daptomycin, was prepared via a semi-synthesis route from daptomycin. The CB-131547 was a pale yellow  
15 amorphous powder, with a solubility at 25 °C of ~80 mg/mL in normal saline.

CB-131547 (60 mg, ~90% pure) is dissolved in 2.5 mL water. The CB-131547 solution is sequentially mixed in order with 5.0 mL methanol, 0.2 mL 1 M calcium acetate (pH 6.0), 2.5 mL propylene glycol, and 1.0 mL 50% (w/v) PEG 4000 to give a final volume of 11.2 mL. The solution is allowed to sit for 4 to 24 hours at 4 °C. CB-  
20 131547 crystals are formed at a yield of ~ 70% with a purity ~ 98.0% as determined by HPLC.

#### EXAMPLE 18

CB-131547 (see Figure x), a cyclic lipopeptide analog of daptomycin, was prepared via a semi-synthesis route from daptomycin. The CB-131547 was a pale yellow  
25 amorphous powder, with a solubility at 25 °C of ~80 mg/mL in normal saline.

CB-131547 (60 mg, ~90% pure) is dissolved in 2.5 mL water. 0.2 mL 1 M calcium acetate (pH 6.0) and 8 mL of isopropanol is added. The solution is allowed to  
30 equilibrate at room temperature (25 °C) for 5 minutes. One mL aliquots of isopropanol are slowly added until the solution becomes cloudy. The solution is stored at room temperature overnight to form crystals.

All publications and patent applications cited in this specification are herein incorporated by reference as if each individual publication or patent application were specifically and individually indicated to be incorporated by reference. Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it will be readily apparent to those of ordinary skill in the art in light of the teachings of this invention that certain changes and modifications may be made thereto without departing from the spirit or scope of the appended claims.

CLAIMS

1. A crystalline or crystal-like lipopeptide or a salt thereof, wherein the lipopeptide is selected from the group consisting of daptomycin, A54145 and a daptomycin-related lipopeptide.

5

2. The crystalline or crystal-like lipopeptide or salt thereof, according to claim 1, wherein the salt is a divalent calcium salt.

10

3. The crystalline or crystal-like lipopeptide antibiotic or salt thereof, according to claim 1, wherein the lipopeptide antibiotic is daptomycin.

4. The crystalline or crystal-like lipopeptide or salt thereof, according to claim 3, wherein the salt is a divalent calcium salt.

15

5. The crystalline daptomycin according to claim 3, wherein an x-ray diffraction pattern of the crystalline daptomycin, using a Cu ( $\lambda=1.54 \text{ \AA}$ ) x-ray source, has a diffraction angle ( $2\theta$ ) = 10.9, 19.2 and 23.3 (degree) or a diffraction angle ( $2\theta$ ) = 19.2, 23.2, 23.4 and 23.6 (degree).

20

6. The crystal-like lipopeptide according to claims 1-4, wherein crystal-like means the compound has crystalline characteristics by birefringence, but does not have crystalline characteristics by x-ray powder diffraction.

25

7. The crystalline or crystal-like daptomycin according to claim 3, wherein the crystalline or crystal-like daptomycin comprises urchin-like or a cluster of needles, needle-like, or rod-like crystals.

30

8. The crystalline or crystal-like daptomycin according to claim 3, wherein the crystalline daptomycin has a purity of at least 95%.

9. The crystalline or crystal-like daptomycin according to claim 3, wherein the crystalline daptomycin or salt thereof has a purity of at least 97%.

10. The crystalline or crystal-like daptomycin according to claim 3, wherein the crystalline daptomycin or salt thereof contains no single impurity greater than 1%.

5 11. The crystalline or crystal-like daptomycin according to any one of claims 8-10, wherein the purity is measured by HPLC.

12. The crystalline or crystal-like lipopeptide according to claim 1, wherein a crystal of the lipopeptide is at least 5  $\mu\text{m}$ .

10 13. The crystalline or crystal-like lipopeptide according to claim 12, wherein the crystal is at least 50  $\mu\text{m}$ .

14. The crystalline or crystal-like lipopeptide according to either of  
15 claims 12 or 13, wherein the lipopeptide is daptomycin.

15. The crystalline lipopeptide according to claim 1, wherein the crystalline lipopeptide exhibits a higher stability than an amorphous form of the lipopeptide.

20 16. The crystalline lipopeptide according to claim 15, wherein the crystalline lipopeptide exhibits higher stability to heat, light, degradation or humidity than the amorphous form.

25 17. The crystalline lipopeptide according to claim 16, wherein the stability is measured by antibiotic activity or degradation of the lipopeptide antibiotic.

30 18. The crystalline lipopeptide according to claim 15, wherein the lipopeptide is daptomycin.

19. The crystalline daptomycin according to claim 18, wherein the crystalline lipopeptide exhibits lower conversion to anhydro-daptomycin or the  $\beta$ -isomer of daptomycin than the amorphous form of daptomycin.

5                   20. The crystalline lipopeptide according to claim 1, which is a daptomycin-related lipopeptide.

21. A pharmaceutical composition comprising a crystalline or crystal-like lipopeptide antibiotic and a pharmaceutically acceptable carrier, wherein the lipopeptide antibiotic is selected from the group consisting of daptomycin,  
10                   A54145, and a daptomycin-related lipopeptide.

22. The pharmaceutical composition according to claim 21, wherein the crystalline or crystal-like lipopeptide is daptomycin.  
15

23. The pharmaceutical composition according to claim 22, wherein the crystalline or crystal-like daptomycin is enterically coated for oral administration.

20                   24. The pharmaceutical composition according to claim 22, wherein the crystalline or crystal-like daptomycin is formulated in a dose of 3 to 75 mg/kg.

25                   25. The pharmaceutical composition according to claim 22, wherein the carrier enhances the oral availability of daptomycin.

26. The pharmaceutical composition according to claim 22, which is in the form of micronized particles or microspheres.

27. A container comprising the pharmaceutical composition according to claim 21.  
30

28. The pharmaceutical composition according to claim 26, which is used as an aerosol.



29. A formulation comprising a crystalline or crystal-like lipopeptide antibiotic and a pharmaceutically acceptable carrier, wherein the lipopeptide antibiotic is selected from the group consisting of daptomycin, A54145 and a daptomycin-related lipopeptide.

30. The formulation according to claim 29, which is a pharmaceutical formulation, a food formulation, a feed formulation, a veterinary formulation, a cosmetic formulation or a personal care formulation.

31. The formulation according to claim 29, that is a pharmaceutical formulation, wherein the formulation further comprises another antibiotic, a stabilizing agent, an agent to aid in absorption, a pH buffering agent or an inorganic salt.

32. The formulation according to claim 29 that is a feed formulation, wherein the formulation further comprises animal feed and may optionally comprise another antibiotic or vitamins.

33. The formulation according to claim 29, that is a personal care formulation, wherein the personal care formulation is a washing formulation, a soap, a shampoo or an antiperspirant.

34. The formulation according to claim 29, that is a veterinary formulation, wherein the formulation is a soap, a shampoo or a pharmaceutical composition.

35. A method for administering a crystalline or crystal-like lipopeptide, a pharmaceutically acceptable salt thereof or a pharmaceutical composition thereof, wherein the lipopeptide is selected from the group consisting of daptomycin, A54145 and a daptomycin-related lipopeptide, comprising the step of administering to a patient in need thereof a therapeutically effective amount of the

crystalline or crystal-like lipopeptide, the pharmaceutically acceptable salt thereof or the pharmaceutical composition thereof.

5           36.    The method according to claim 35, wherein the lipopeptide antibiotic has a purity of greater than 95%.

          37.    The method according to claim 36, wherein the lipopeptide antibiotic is daptomycin.

10           38.    The method according to claim 37, wherein the daptomycin is a crystalline daptomycin and an x-ray diffraction pattern of the crystalline daptomycin has a diffraction angle ( $2\theta$ ) = 10.9, 19.2 and 23.3 (degree) using a Cu ( $\lambda=1.54 \text{ \AA}$ ) x-ray source.

15           39.    The method according to claim 37, wherein the daptomycin is a crystal-like daptomycin and the crystal-like daptomycin has crystalline characteristics by birefringence but does not have crystalline characteristics by x-ray powder diffraction.

20           40.    The method according to claim 35, wherein the crystalline or crystal-like lipopeptide is administered as a micronized particle.

          41.    The method according to claim 35, wherein the crystalline or crystal-like lipopeptide is administered as a targeted release form.

25           42.    The method according to either of claims 40 or 41, wherein the lipopeptide is daptomycin.

          43.    The method according to claim 35, wherein the oral administration  
30 is done subcutaneously, intravenously or intramuscularly.

44. A method for storing a lipopeptide, wherein the lipopeptide is selected from the group consisting of daptomycin, A54145 and a daptomycin-related lipopeptide, comprising the steps of

- a) providing a dissolved solution of a lipopeptide;
- b) crystallizing or precipitating the lipopeptide;
- c) collecting and drying the lipopeptide; and
- d) storing the lipopeptide;

wherein the crystalline or crystal-like lipopeptide is more stable than an amorphous form of the lipopeptide.

45. A method for manufacturing a crystalline or crystal-like lipopeptide, wherein the lipopeptide is selected from the group consisting of daptomycin, A54145 and a daptomycin-related lipopeptide, comprising the steps of

- a) providing an amorphous form of a lipopeptide;
- b) crystallizing or precipitating the lipopeptide; and
- c) collecting the crystalline or crystal-like lipopeptide.

46. The method according to claim 44, wherein said collecting is performed by filtration.

47. The method according to claim 46, further comprising the step of washing the lipopeptide after step b).

48. The method according to either of claims 45 or 46, further comprising the step of drying the lipopeptide after step c).

49. The method according to claim 48, further comprising the step of sterilizing the lipopeptide after drying.

50. The method according to claim 45, wherein step c) is performed under sterile conditions.

51. The method according to claim 50, wherein step b) is performed under sterile conditions.

52. The method according to claim 51, further comprising the step of drying the lipopeptide after step c) under sterile conditions.

5

53. The method according to claim 44, wherein the purity of the crystalline lipopeptide is higher than the amorphous form of the lipopeptide.

54. The method according to claim 53, wherein the purity of the amorphous form is approximately 90% and the purity of the crystalline or crystal-like form is greater than 95%.

10

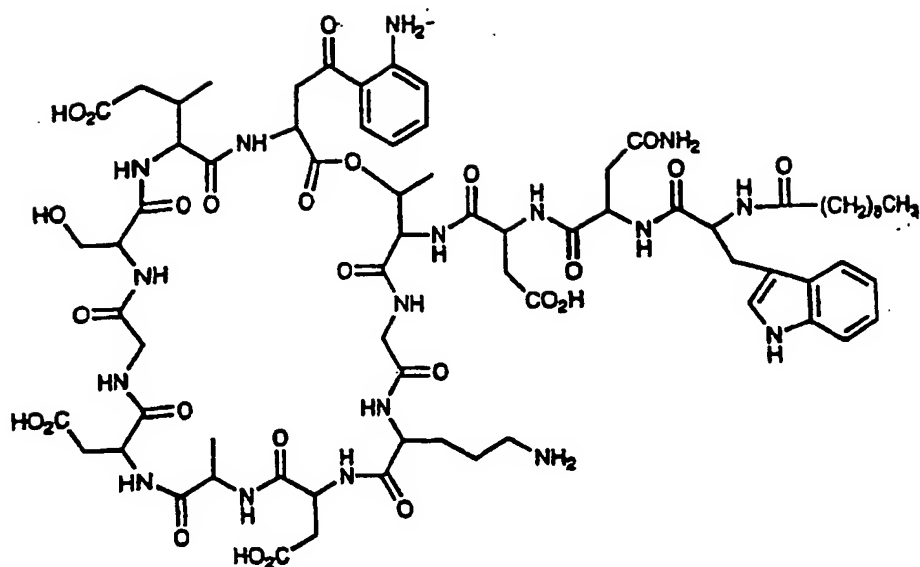
55. The method according to claim 53, wherein the purity of the amorphous form is approximately 93%, and the purity of the crystalline or crystal-like form is greater than 95%.

15

56. The method according to claim 53, wherein the purity of the amorphous form is approximately 93%, and the purity of the crystalline or crystal-like form is approximately 98%.

20

### Figure 1



**Figure 2**

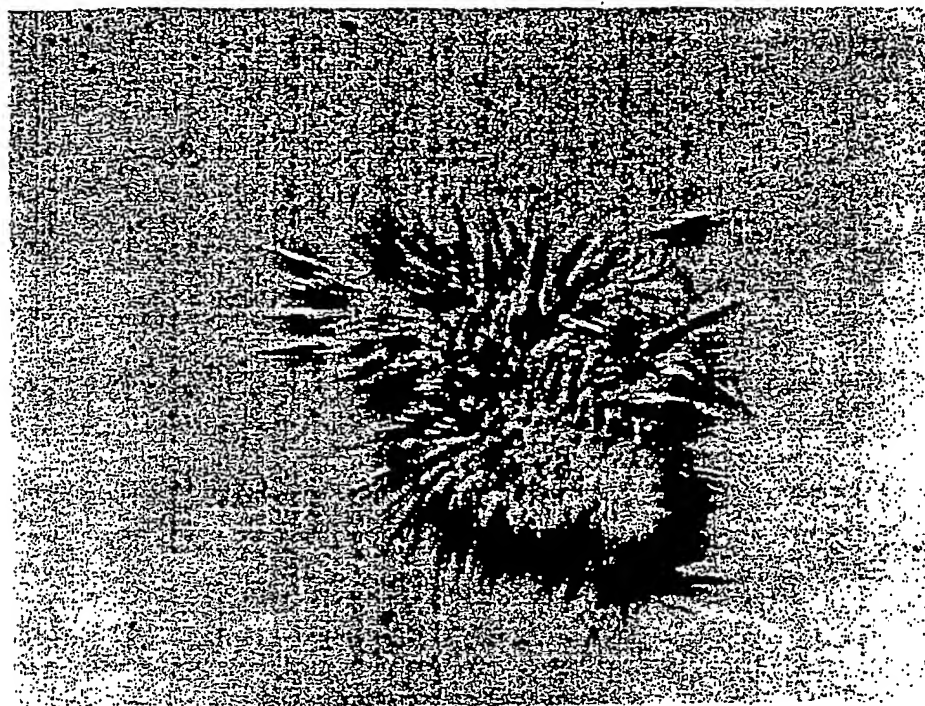


Figure 3

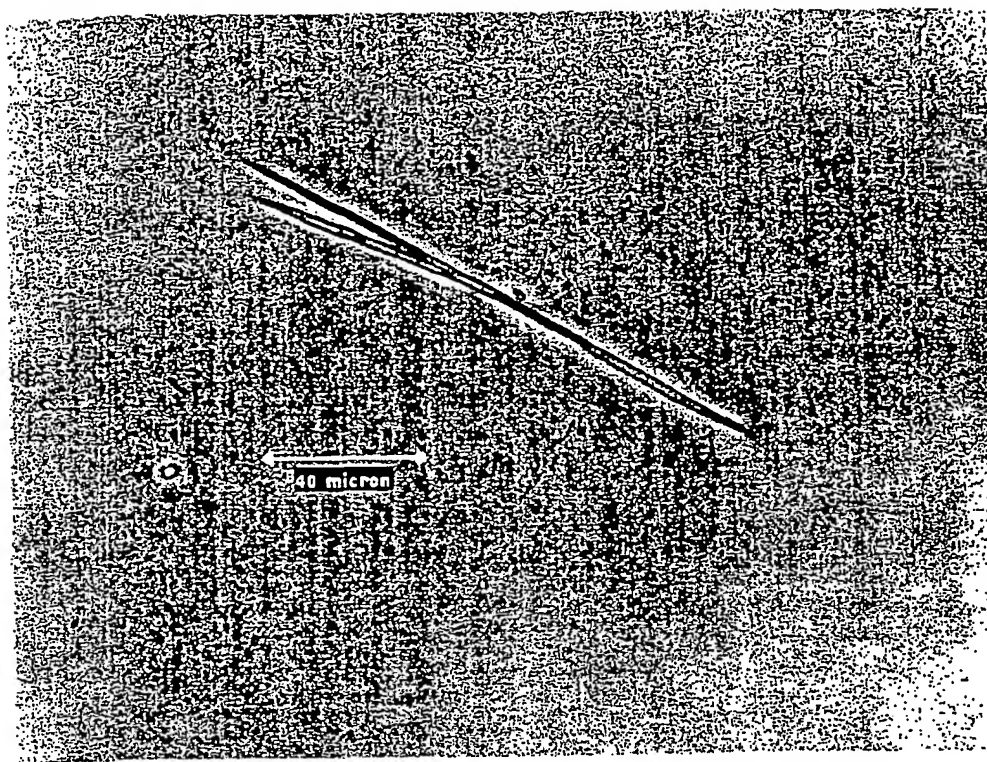


Figure 4

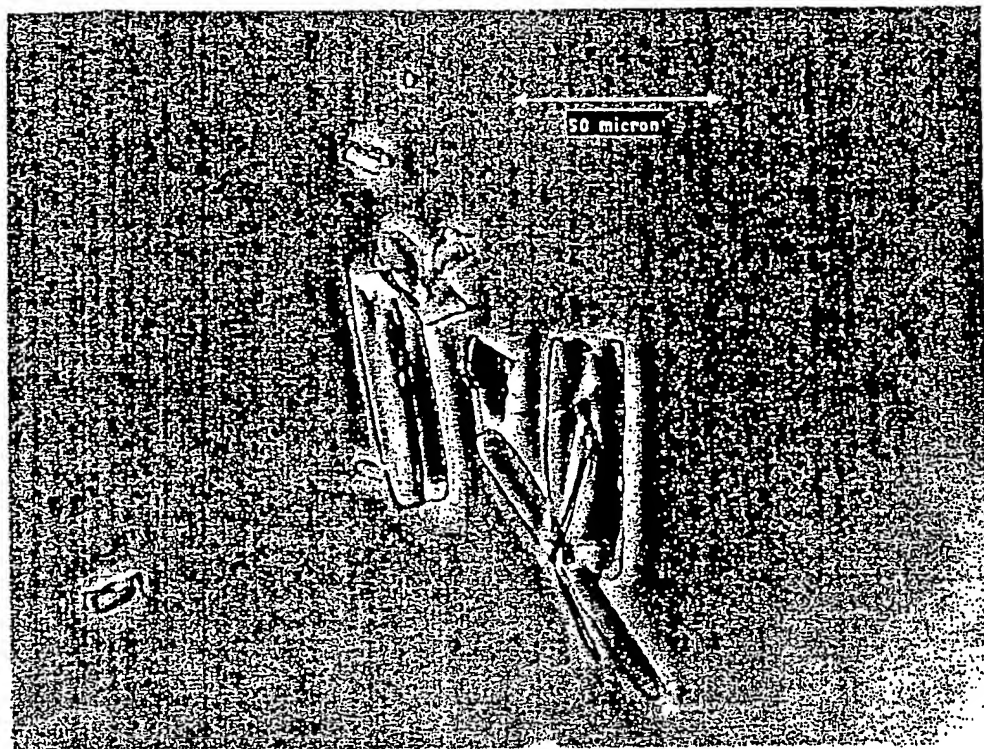




Figure 5

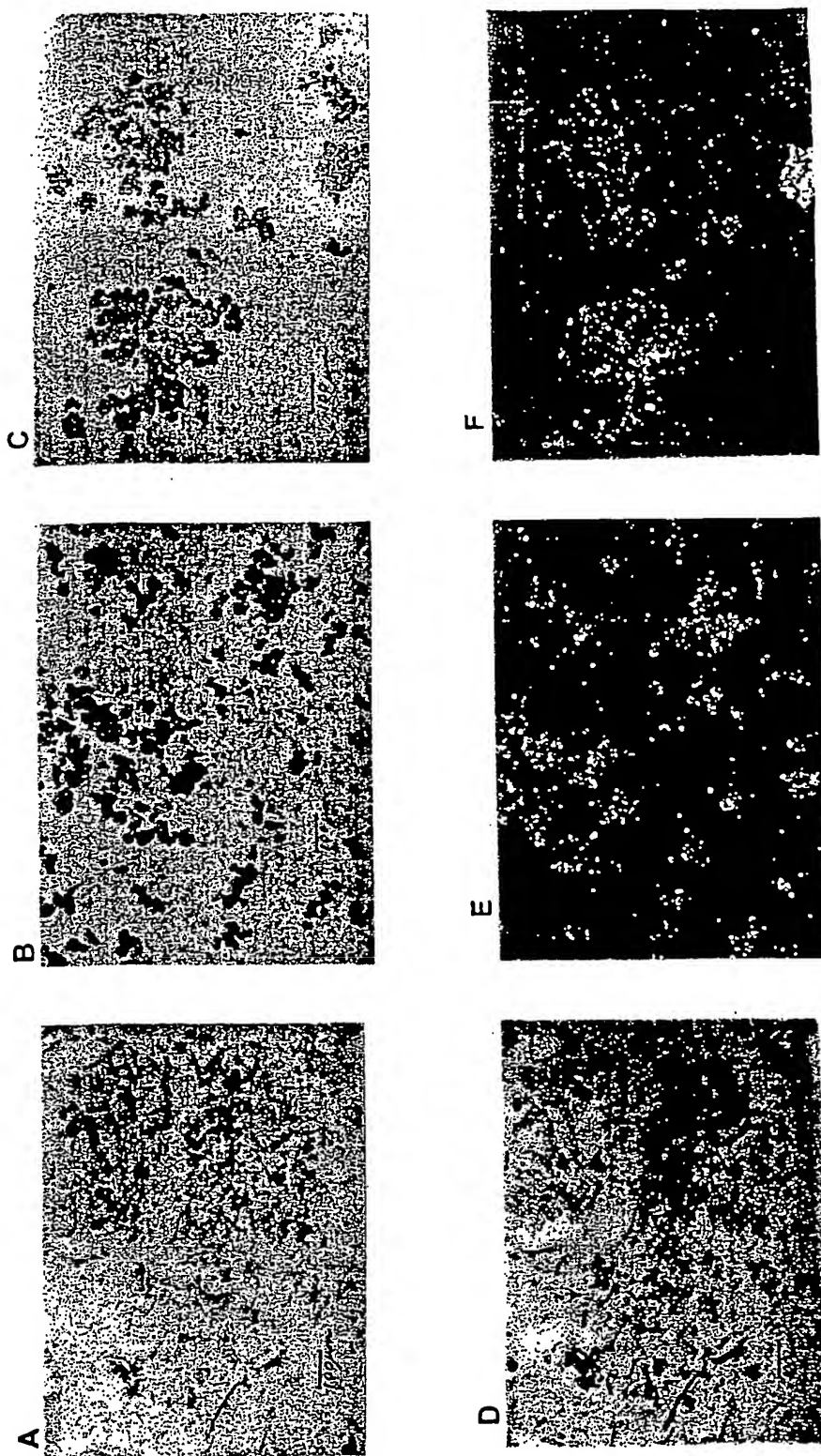


Figure 6

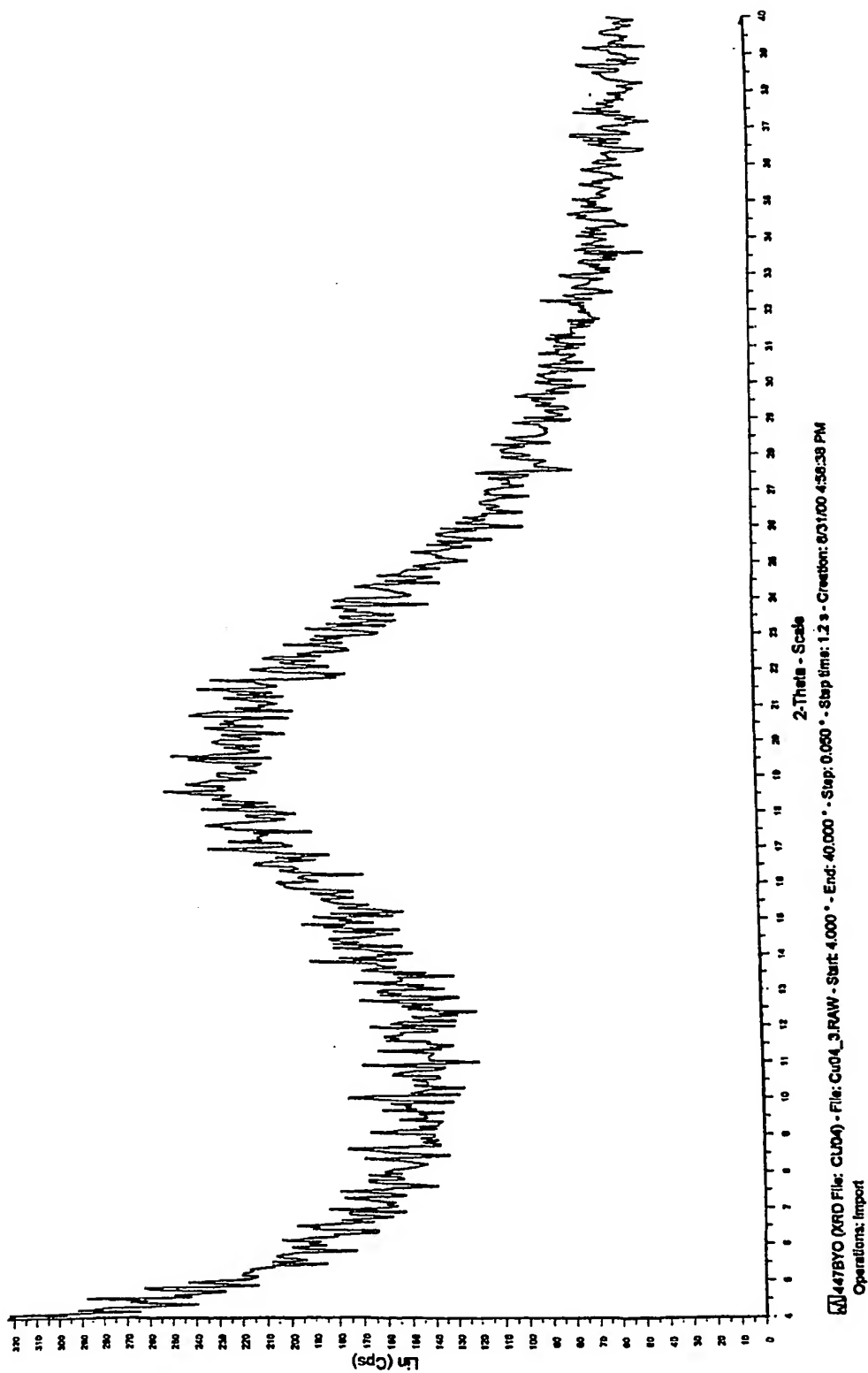


Figure 7

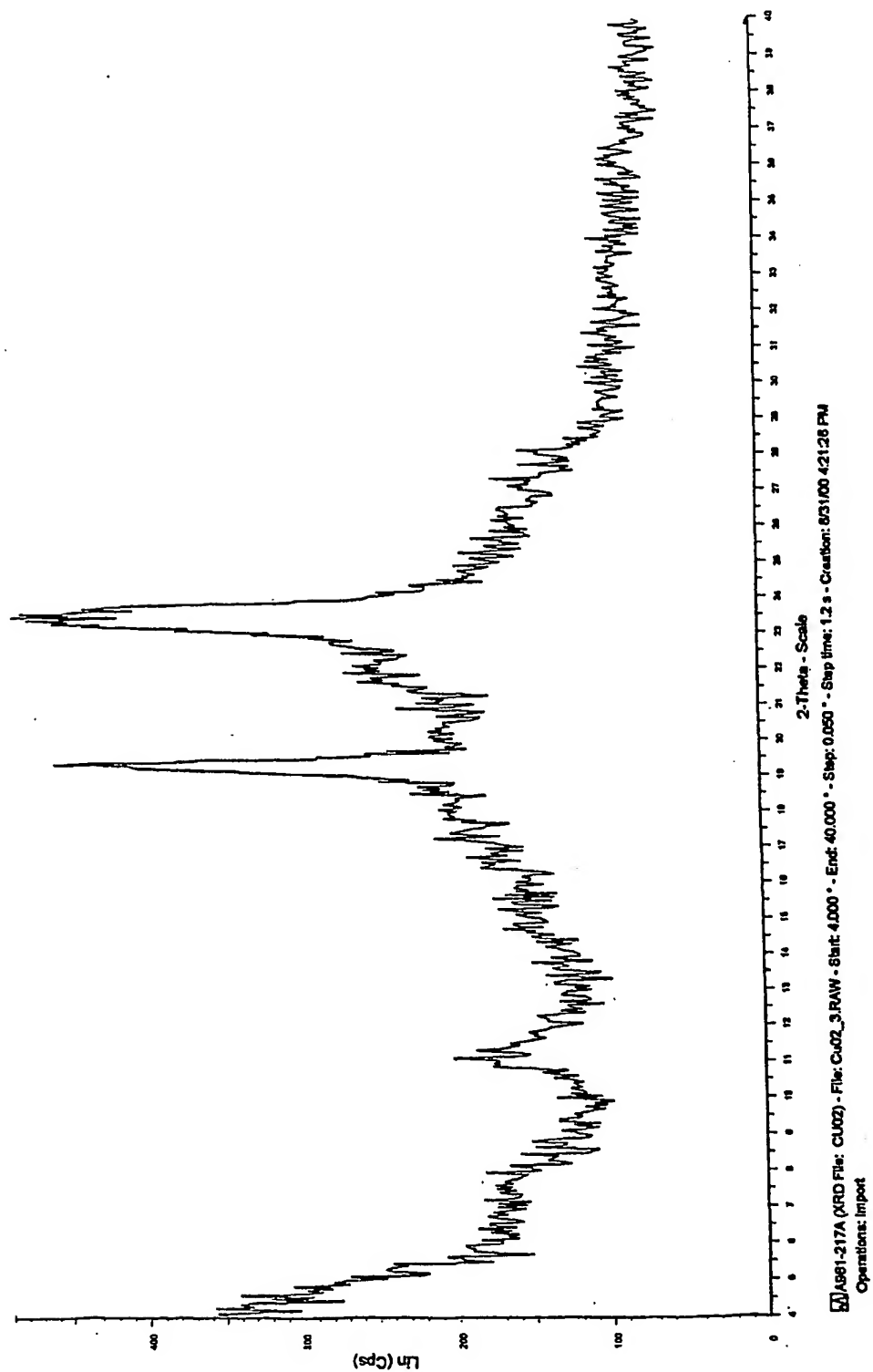


Figure 8

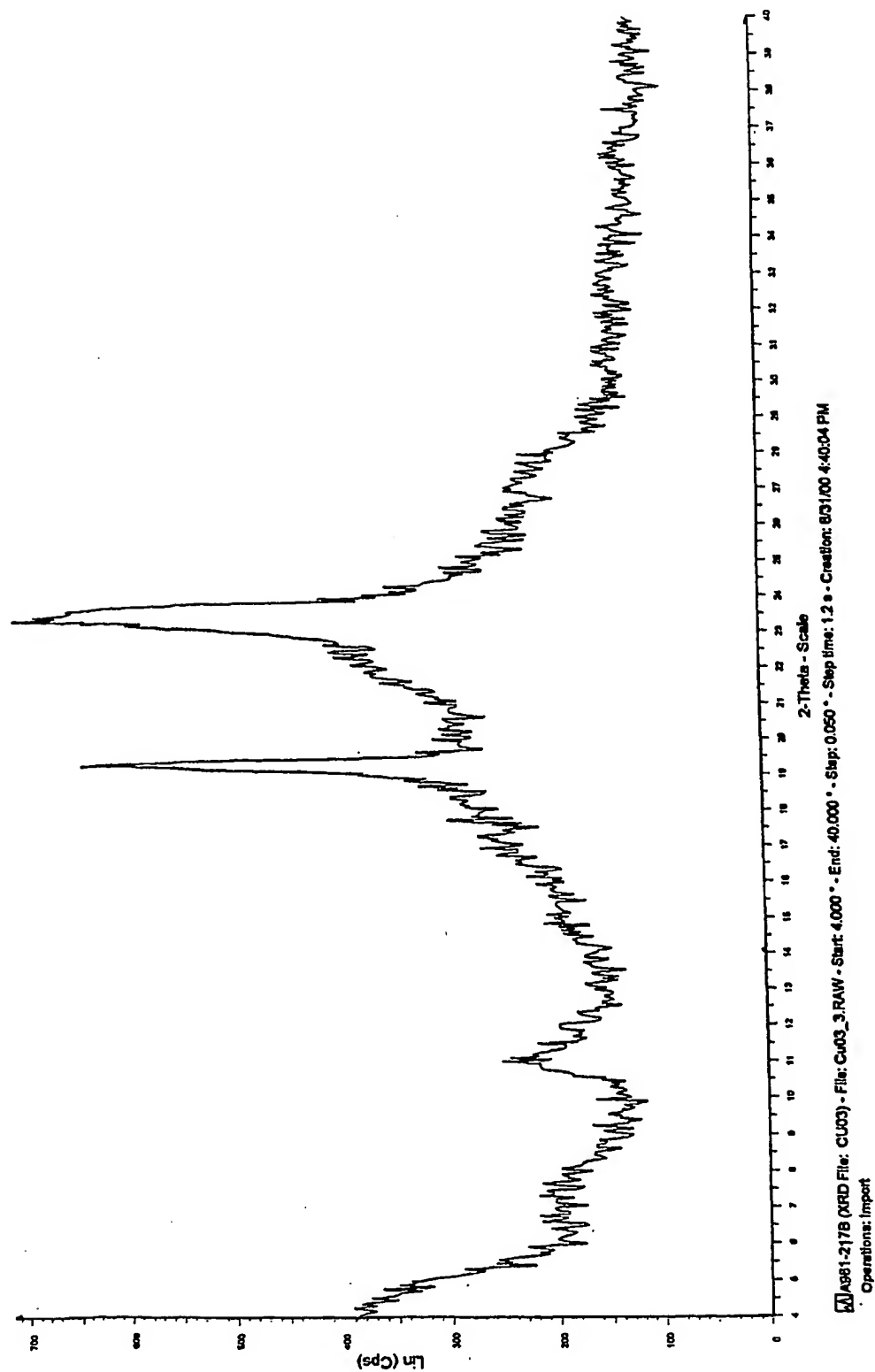


Figure 9

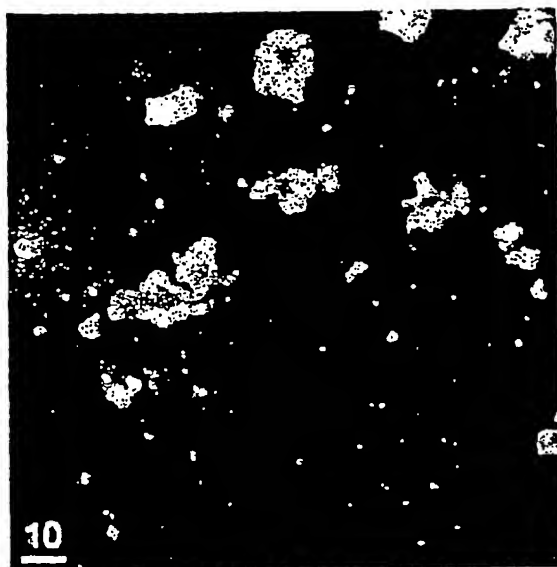


Figure 10

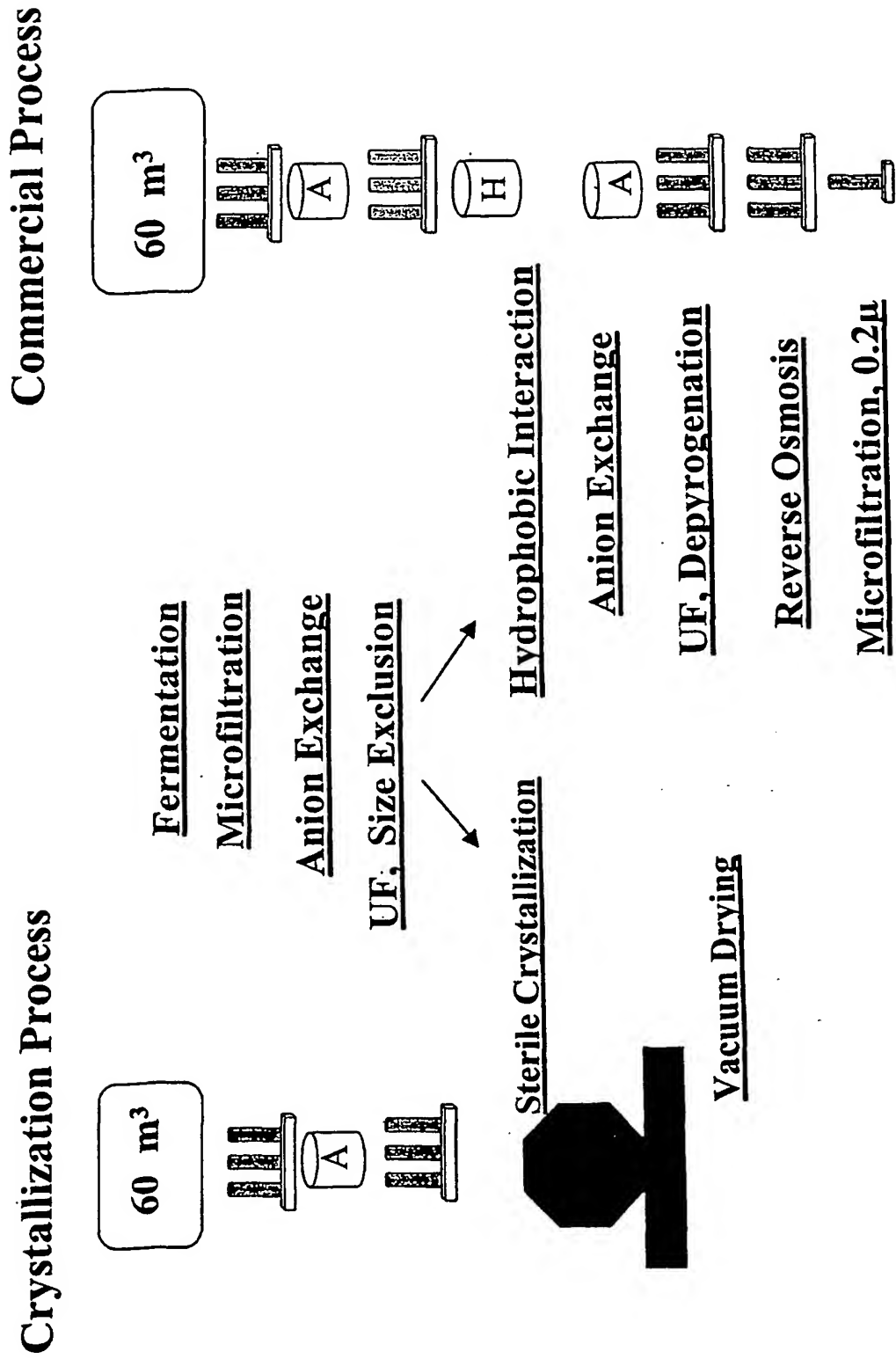


Figure 11

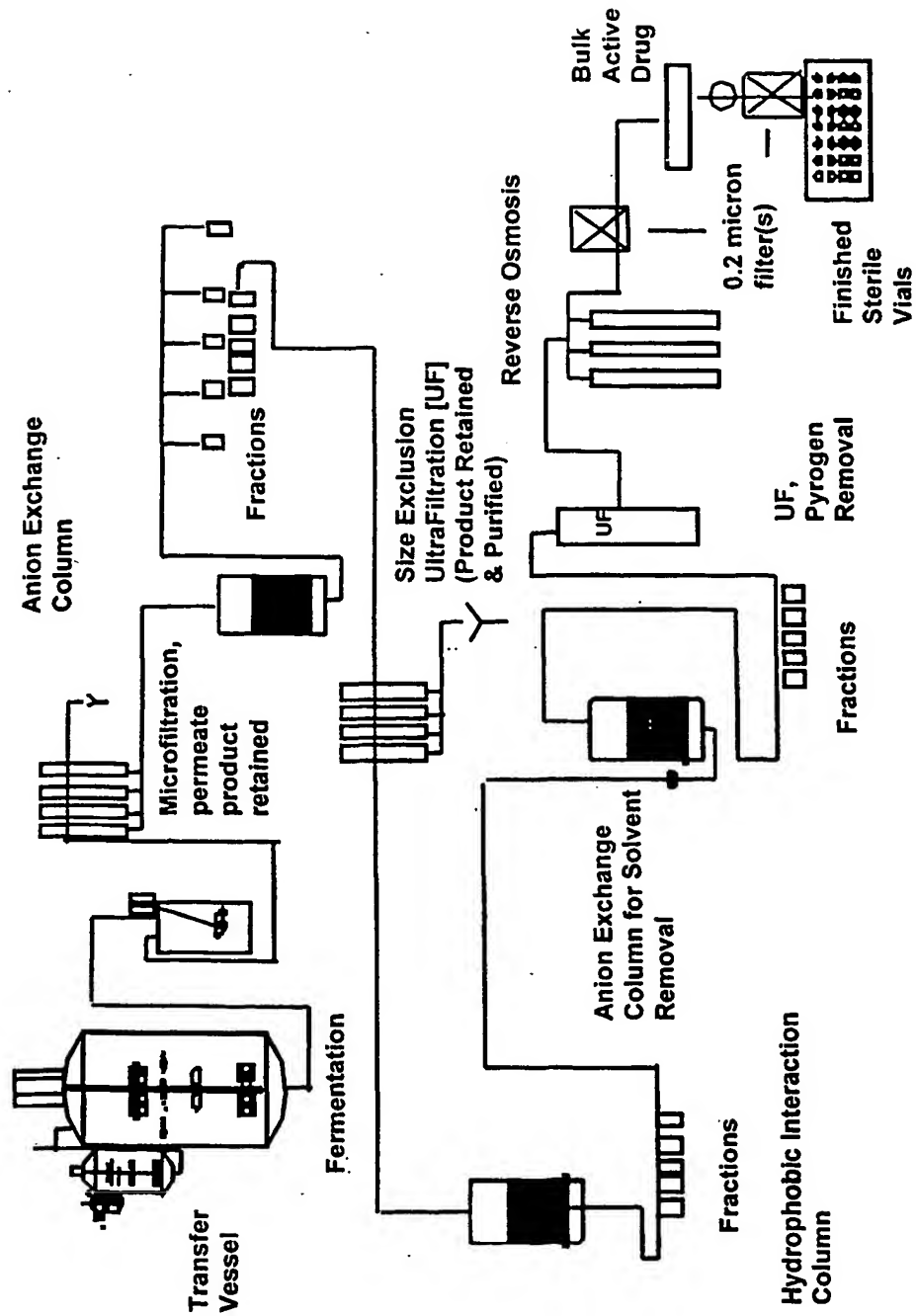


Figure 12

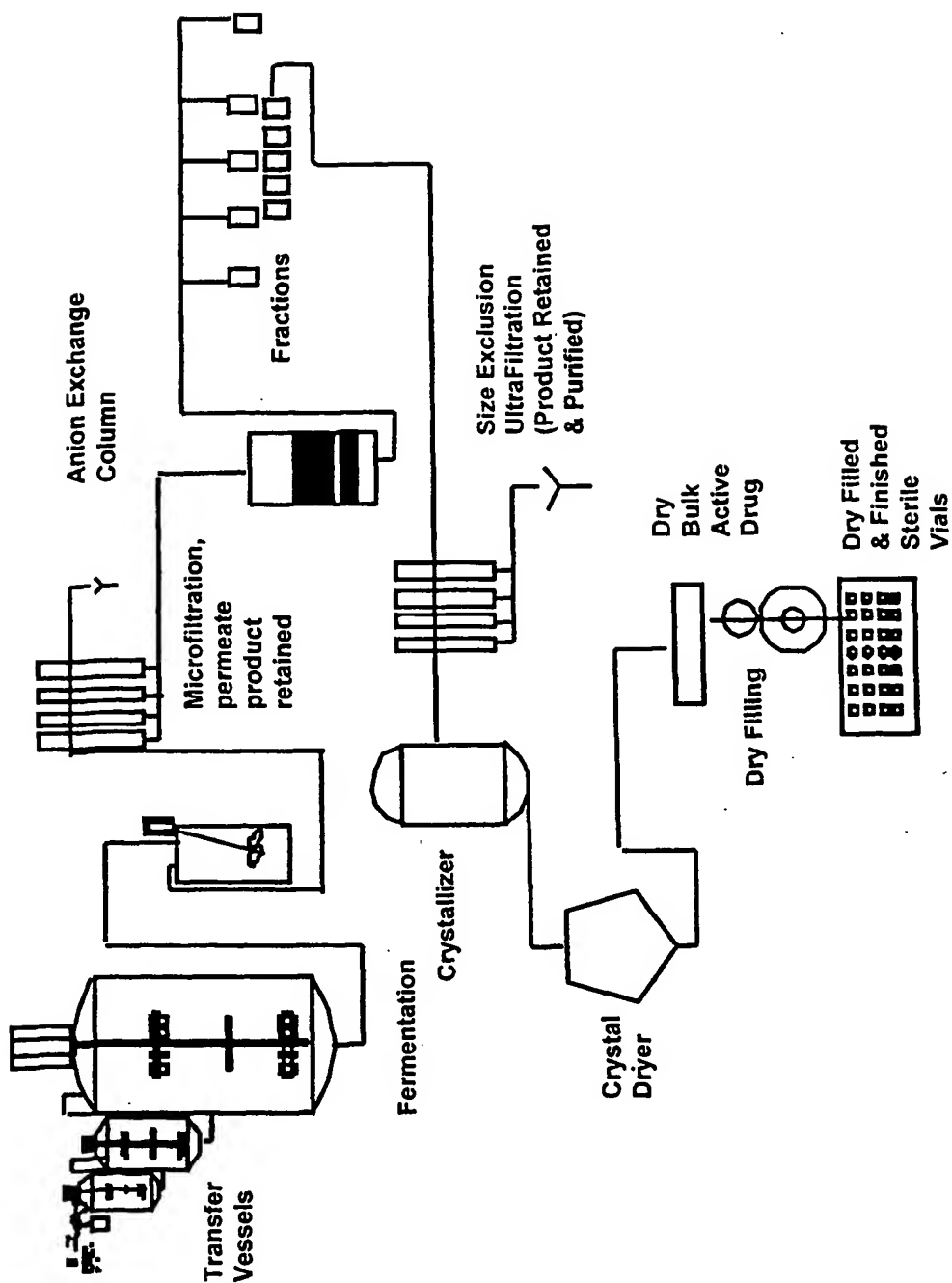




Figure 13

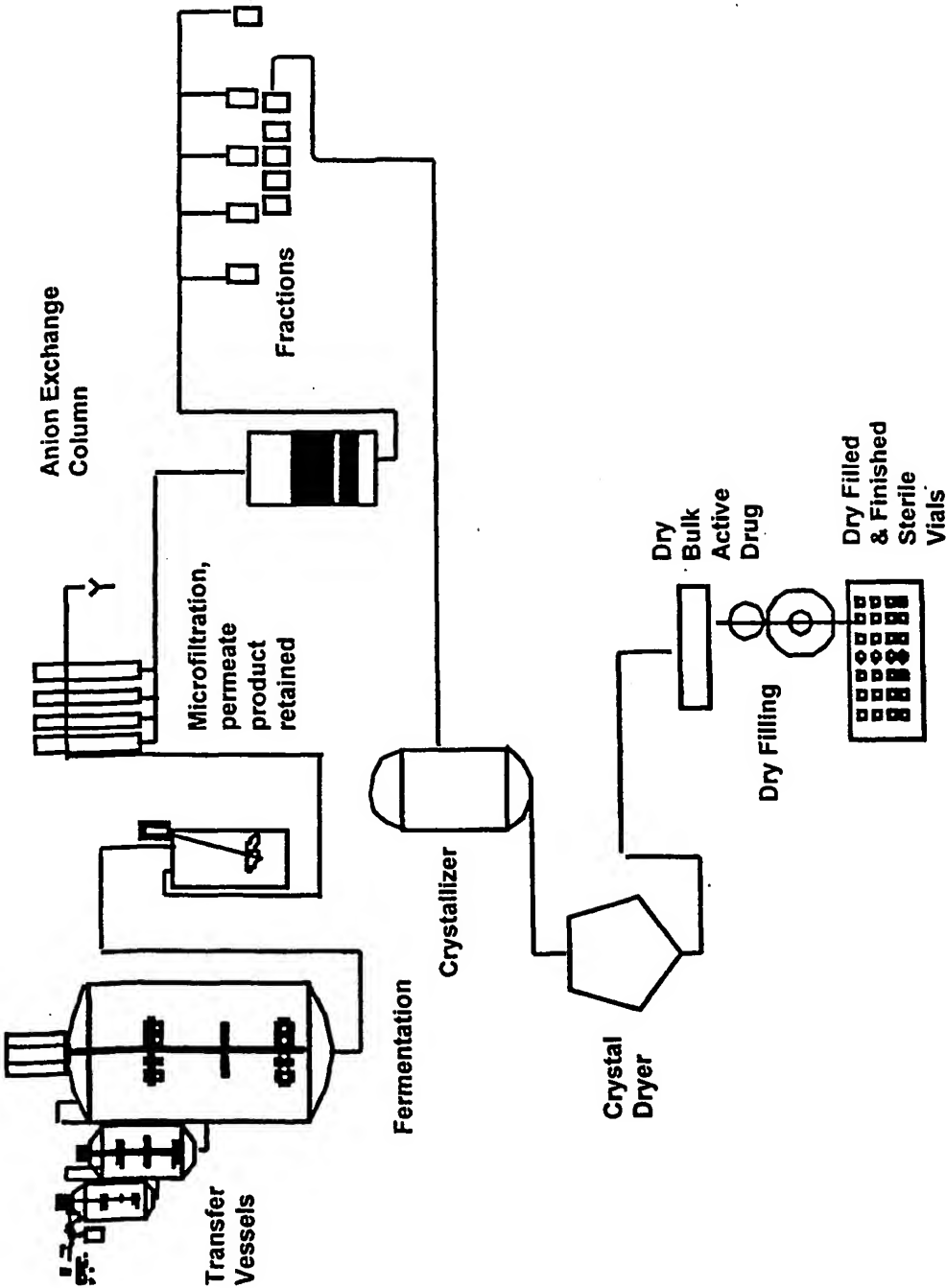


Figure 14

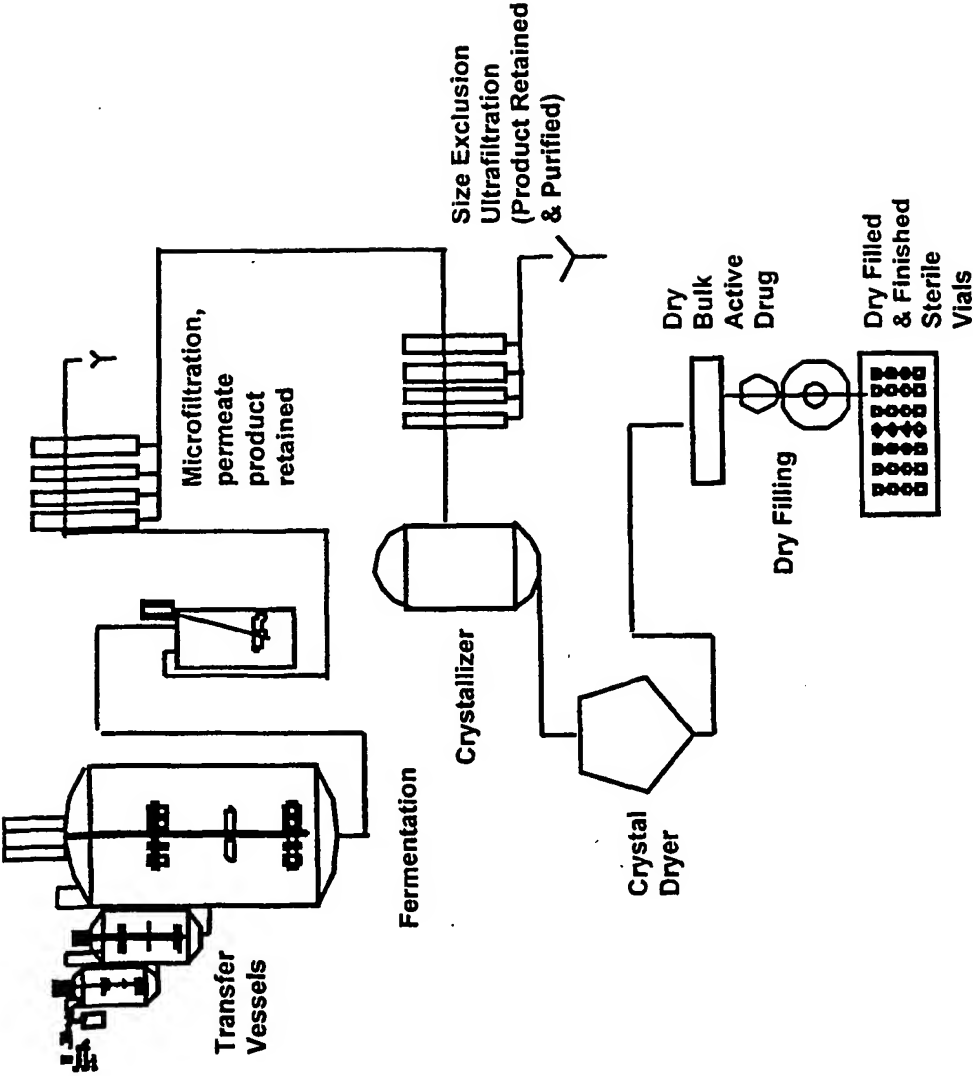
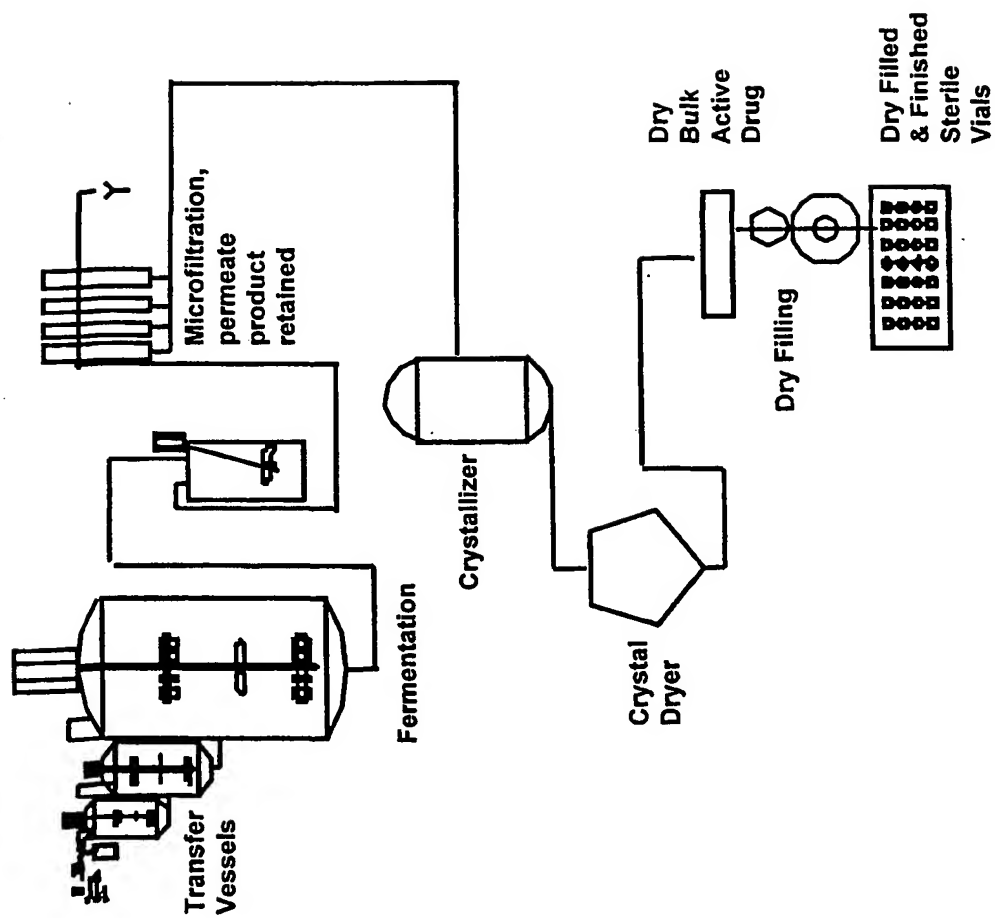
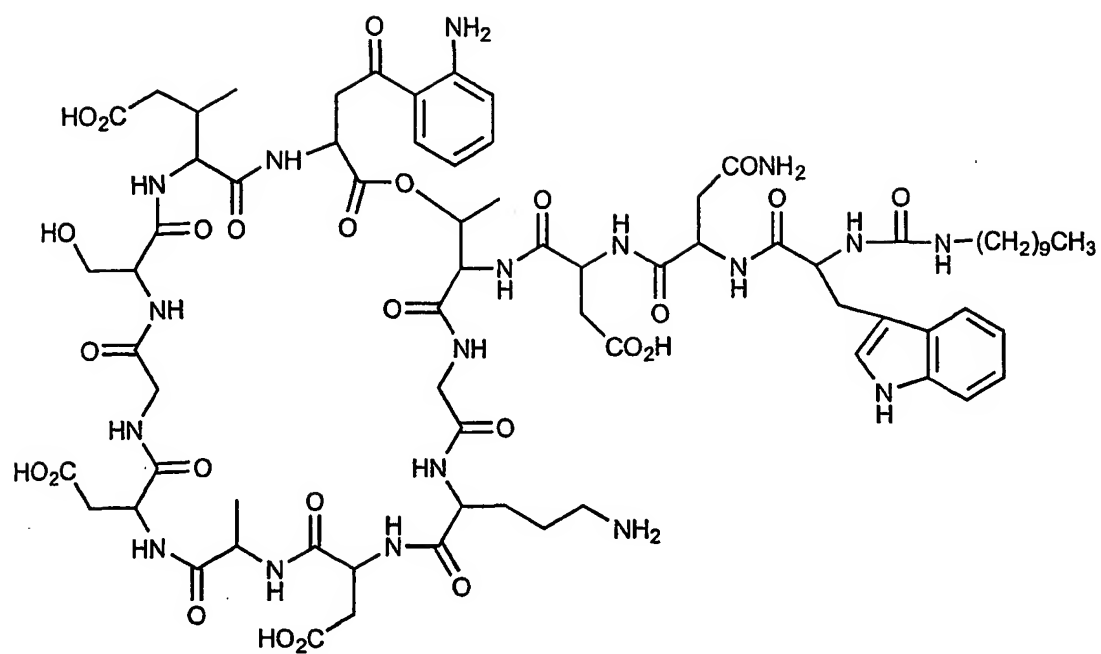


Figure 15



Figur 16



## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/48886

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(7) : C07K 7/64, 14/36

US CL : 514/9, 11; 530/317, 321

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : NONE

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS ONLINE, MEDLINE, WPIDS, EAST, WEST

search terms: crystal, crystallization, daptomycin, A54146, A-21978C

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 386 951 A2 (ELI LILLY AND COMPANY) 09 December 1990, see entire document.	1-5, 7-56
A	US 4,537,717 A (ABBOT et al.) 27 August 1985, see entire document.	1-5, 7-56
A	US 5,039,789 A (FUKUDA et al.) 13 August 1991, see entire document.	1-5, 7-56
A	US 4,994,270 A (BOECK et al.) 19 February 1991, see entire document.	1-5, 7-56



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

26 JUNE 2002

Date of mailing of the international search report

11 JUL 2002

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# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/48886

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claims Nos.: 6  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.  
☐ No protest accompanied the payment of additional search fees.